



(12)

## EUROPEAN PATENT APPLICATION

(21) Application number: **91302521.9**

(51) Int. Cl.<sup>5</sup>: **A61B 5/14, A61B 5/00**

(22) Date of filing: **22.03.91**

(30) Priority: **26.03.90 US 499085**

(43) Date of publication of application:  
**02.10.91 Bulletin 91/40**

(84) Designated Contracting States:  
**AT BE CH DE DK ES FR GB GR IT LI LU NL SE**

(71) Applicant: **CASCADE MEDICAL, INC.**  
10180 Viking Drive  
Eden Prairie, Minnesota 55344 (US)

(72) Inventor: **Anderson, Paul J.**  
8243 Thomas Avenue South  
Bloomington, Minnesota 55431 (US)

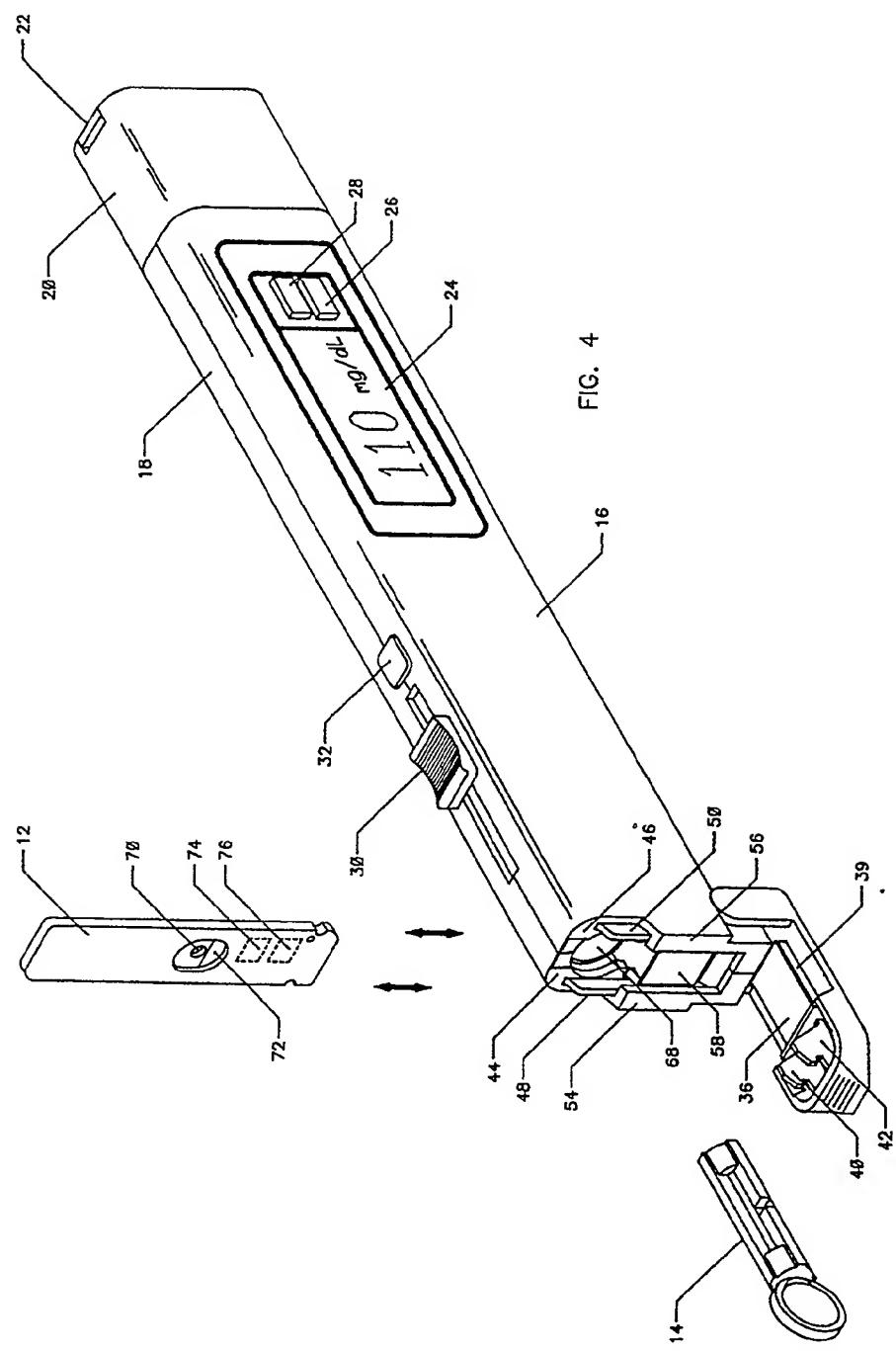
Inventor: **Jessen, Ross A.**  
305 Waters Edge Drive  
Chaska, Minnesota 55318 (US)  
Inventor: **Linde, David R.**  
6537 Meadowlark Lane  
Maple Grove, Minnesota 55369 (US)  
Inventor: **Jones, Richard E.**  
7901 Rhode Island Circle  
Bloomington, Minnesota 55438 (US)  
Inventor: **Walter, Bert**  
809 Park Place Drive  
Mendota Heights, Minnesota 55118 (US)  
Inventor: **Christenson, Paul E.**  
6886 133rd Street West  
Apple Valley, Minnesota 55124 (US)

(74) Representative: **Parr, Ronald Edward R.E. Parr & Co.**  
Colman House Station Road  
Knowle Solihull West Midlands B93 0HL (GB)

(54) **Medical diagnostic system.**

(57) Hand-held shirt-pocket portable instrument (10) for quantitative measurement of glucose or analysis in biological fluids. The system (10) accepts a disposable diagnostic reagent test device (12) which has a reagent carrier and a mode of identifying to the instrument the characteristics of the chemical reagent used. The instrument includes a housing structure having a visual LCD readout (24), a microprocessor (249), and photosensing circuitry (207) which measures the change of color of the reagent carrier upon reaction of the reagent in the disposable test device (12).

The housing also includes a spring arrangement (132,134) for actuating a disposable lancet (14) into the skin for generating blood. The disposable diagnostic reagent unit (12) includes a configuration for transporting the blood into the reagent unit (12). The system includes verification and calibration sequences for the electronics, the chemical reagent of an unused disposable unit, the presence of a blood sample, and the ambient temperature. The system also provides for storing a plurality of analysis readings.



**1. Field of the invention -**

The present invention pertains to a system for use in the analysis of a liquid and to a process for measuring a component quality or quantity of a liquid. The invention relates particularly, but not exclusively, to the medical field, for example to a medical monitoring diagnostic system for sampling and analyzing blood or any components of blood for specific readings as to qualities of the blood. One specific use of the present invention is for sensing the accumulating of blood glucose in diabetics. In a preferred form the invention relates to a portable, pocket-size, battery-operated, diagnostic system for use with a disposable diagnostic reagent unit. In one form of the invention the system is used with a disposable lancet.

A test device or reagent unit suitable for use with a system of the present invention is described and claimed in a European application filed on the same day as the present application and by the same applicant, under the title "Disposable Reagent Unit". That European application claims priority from U.S. Patent Application 07/499 187 filed 26 March 1990.

The invention is described hereinafter with particular reference to the analysis of blood with reference to its glucose content but applies, mutatis mutandis, to the analysis of analytes in liquids generally.

**2. Description of the Prior Art -**

Prior art blood glucose devices have operated on the principle of taking blood from an individual by a variety of methods, such as by a separate needle or lancing device. An individual then had to coat a separate unit carrying chemistry with the blood, time the chemical reaction for about 60 seconds, wipe or remove the blood sample from the unit, and insert the blood-coated unit into a blood glucose meter or make a visual personal comparison against a color standard.

There are numerous blood glucose meters in the market-place, most of them bulky and not easily pocketable. The instruments usually have to be carried in a large handbag, or an individual's briefcase, or left at home such as in the bathroom or the bedroom, or on a counter or a table.

The prior art medical apparatuses for sensing blood glucose required that an individual have separately available a needle or lance for extracting blood from the individual, units carrying blood chemistry for creating a chemical reaction with respect to the blood glucose and changing color, and a blood glucose meter for reading the change in color indicating the blood glucose level. The level of blood glucose, when measured by a glucometer, is read from a unit carrying the blood chemistry through the well known process of reflectometers based on the principle of

glucose oxidation.

Some of the monitor/reagent unit systems that are now available on the market have multiple sequential steps that the patient must follow at exact time intervals. Each step is subject to error by the patient. As in most monitors, it is the patient's responsibility to periodically calibrate the monitor against known color standards; validate the efficacy of the reagent units and technique by immersing the units in a control solution of known glucose content; and, then comparing the color change visually against the color standard or by using a calibrated monitor. These types of prior art systems are subject of course to human error.

The procedure for obtaining accurate results from the time a drop of blood is placed on a reagent unit pad to the time the pad color change may be read in the glucose monitor is as follows. The patient must stick himself/herself with a lancet. A drop of blood must be squeezed to the surface of the skin. The blood must then be carefully placed on the reagent pad, making sure to cover the pad completely and that the pad is never touched by the finger of the patient to prevent contamination. Once the sample has been applied to the surface of the reagent pad, the patient must press a timer on the monitor. At the end of the timing, the patient must wipe, blot or wash the unit off, using a careful technique. And for most units, the patient must place the reacted reagent unit into the monitor, and press a test button or close a hatch to obtain results. Prior art commercially available comparable reagent units or monitors require operator intervention in a prescribed sequence at exact time intervals. The prior art monitors are subject to operator error, sequence errors, timing errors, and technique errors.

The prior art reagent units are also subject to contamination which may affect accuracy of measurement.

A representative patent is U.S. Patent No. 4,787,398, entitled "Glucose Medical Monitoring System", issued on November 29, 1988. In a preferred form, the present invention overcomes the disadvantages of the prior art by providing an integrated handheld pocketable blood glucose monitoring meter which includes an attachable disposable lancet, reagent test device for blood glucose, unit carrying a chemical reagent chemistry, and a wick for, transporting the blood to the blood sensing reagent, resulting in a readout of a level of the blood glucose.

**SUMMARY OF THE INVENTION**

One general purpose of the present invention is a portable, shirt-pocket-size, battery-operated diagnostic system for use by health professionals and/or lay patients for the detection and measurement of certain selected chemical agents or substances for the purpose of diagnosis and/or treatment of disease. The

system application is not restricted to use with human beings as to the sampling of blood glucose. The system may also be extended to veterinary medicine animals, and can also have uses in the agricultural field, such as measurement of glucose in grapes in the wine industry by way of example. One such medical application is for insulin dependent and non-insulin dependent diabetics for the measurement of glucose in serum, plasma, and/or whole blood. the particular quantity to be measured is glucose through the principles of either reflectance, absorption or potentiometric measurement by electronic circuitry although other quantities can be measured.

Another purpose of the present invention is to provide a hand-held pocketable medical measurement system including the engaging of a disposable lancet and a disposable diagnostic reagent unit carrying the blood sensing reagent for sensing readings of the blood, such as blood glucose level. The medical system is cost effective and simple to operate by an individual. The reading, such as an individual's glucose level, is displayed on an LCD display on the side of a housing of the medical system which approximates the size of an ordinary page highlighter which can be carried in an individual's shirt pocket. The disposable diagnostic reagent units in sterile packages and disposable lancets can be carried in a corresponding packets. The housing structure resembling a page highlighter contains the hand-held pocketable medical system. A like housing structure resembling a highlighter carries the extra supply of disposable units. The design of the present invention provides for the utmost peace of mind for the individual.

According to one embodiment of the present invention, there is provided a hand-held pocketable medical system including an electromechanical structure for actuating a disposable lancet about a disposable diagnostic reagent unit which engages onto the system. The disposable diagnostic reagent unit enables a blood sample inside a finger or on the finger surface to be transferred to the blood reagent chemistry. The electromechanical structure includes a spring actuated configuration for movement of a hammer mechanism. The disposable lancet unit and diagnostic reagent unit engage and slide into the end of the hand-held pocketable medical system, and are easily releasable and disposable after a single use. The disposable lancet can be reused as may be required.

The hand-held medical system includes a light tight compartment with photosensing electronics connected to a microprocessor for analyzing the properties of the blood sensing chemistry in the disposable diagnostic reagent unit, and for displaying a readout and storing previous readouts. The electronics includes verification sequences for verifying operability of the electronics including annunciating of a low battery condition, for verifying the condition of a unused disposable unit, for verifying the presence of

a blood sample and for subsequently providing multiple readings to provide for an averaging of results. The microprocessor can be programmed to measure other quantities.

5 According to other embodiments of the present invention, there is provided a disposable diagnostic reagent unit with a transporting action where a wick serves as the transport structure for the blood. There is also provided a disposable lancet unit in the hand held medical device for the piercing of an individual's skin.

One significant aspect and feature of the present invention is a hand-held pocketable diagnostic medical monitoring system which is utilized for extracting 15 a blood sample from the body, subjecting the sample to chemical analysis, and visually displaying the numerical results to the individual. A disposable diagnostic reagent unit carries the blood sensing chemistry consisting of a reagent unit for either delivering blood to the reagent or for causing the reagent to be delivered to the blood. Additional disposable units can be carried in a corresponding structure similar to that of the medical system.

Another significant aspect and feature of the present invention is a housing like structure which is electromechanical, and where a button is pushed for actuating a firing mechanism in the housing structure against the disposable lancet contained therein through the spring driven structure. A hammer return spring returns the firing mechanism back to an original rest position and at about the same time, a return spring removes the sharp point of the lancet from the finger.

35 A further significant aspect and feature of the present invention is a hand-held pocketable diagnostic medical monitoring system which provides blood glucose readings where the disposable diagnostic reagent unit carries glucose-oxidase or like chemical reagent. Once the reagent material undergoes a colorimetric, potentiometric, or absorption action proportional to the blood glucose concentration, the microprocessor circuitry through the reflectance colorimeter provides for subsequent processing of the photosensing of the blood chemistry for displaying of 40 the results on an LCD display.

45 Another significant aspect and feature of the present invention is a system which utilizes a slidable disposable diagnostic reagent unit. The reagent unit is a transport mechanism for transporting a fluid or liquid to the reagent unit.

50 Still another significant aspect and feature of the present invention is a system which inherently through mechanical operation pushes the disposal lance out of the housing to drop into a basket for disposal.

55 Having thus described embodiments of the present invention, it is principal objects hereof to provide a pocketable diagnostic medical monitoring system,

including a disposable lancet and a disposable diagnostic reagent unit which carries blood sensing reagent material and which engages onto the system for providing a subsequent readout on a visual display of the system of a quality of the blood by the system. The system can be broadly extended to a system for measurement of a quantity of a substance in a particular fluid or material, and is not to be construed as strictly limited to medical applications, as the system can be used in industry, commercial, agricultural, consumer or even veterinary environments as examples.

One object of the present invention is to provide a hand-held pocketable diagnostic medical monitoring system with a disposable lancet and a disposable diagnostic reagent unit which engages onto the electromechanical assembly of the medical system. The disposable diagnostic reagent unit carries blood sensing reagent material for sensing components of the blood for qualities such as glucose level. Other qualities of fluid which can be measured are cholesterol, urea, nitrogen, hemoglobin, alcohol, protein or other qualities of the blood with appropriate reagent material.

Another object of the present invention is an electromechanical assembly which contains the microprocessor including the software, mechanical and electromechanical apparatus, batteries, and related circuitry that causes the electrical and electromechanical functional operation. The diagnostic is a disposable reagent unit containing the lancet for obtaining a blood sample, typically from a person's finger or toe, and a chemical impregnated reagent material that reacts with the presence of blood. The chemical reagent is sealed inside the reagent unit housing minimizing the effects of contamination from fingers, moisture, and light, thus improving accuracy and precision of measurement by stabilizing the oxidation reduction or chemical reaction of the reagent prior to use. The sensor in the assembly detects and measures via absorption, potentiometric, or reflectance analysis the amount of glucose or other blood quantity present. This analog data is provided and converted to a digital readout display quantifying glucose in milligrams per deciliter (mg/dl) or MMOL/L.

An additional object of the present invention is a self-contained automatic medical monitoring system. All operations and performance of the system are performed automatically, mechanically and electronically in proper sequences. Accuracy and precision of the measurement is enhanced because errors due to operator interpretation, operator technique, and timing of events, are removed from operator control because of microprocessor based system operation and a lot to lot as well as a test strip calibration.

Still another object of the present invention is a medical diagnostic system which is software controlled and software intelligent. The system is self-calibrating through control commands by the software, and

also based on a lot to lot material in the reagent unit and a calibration square on the inside of the dust cover.

## 5 DESCRIPTION OF THE PREFERRED EMBODIMENTS

There are now described, by way of example and with reference to the accompanying drawings, preferred embodiments of the present invention.

In the drawings:

**FIG. 1** illustrates a perspective view of an embodiment of a glucose medical diagnostic system;

**FIG. 2** illustrates a perspective view of the back-side of the medical diagnostic system;

**FIG. 3** illustrates a perspective view of the medical diagnostic system showing the battery case;

**FIG. 4** illustrates a perspective view of the medical diagnostic system ready to receive a disposable lancet and a disposable diagnostic reagent strip;

**FIG. 5** illustrates an exploded view in perspective of the medical diagnostic system;

**FIG. 6** illustrates an exploded view in perspective of the optics head;

**FIG. 7** illustrates an exploded view in perspective of the firing mechanism assembly;

**FIG. 8** illustrates a perspective view of the dust cover;

**FIG. 9** illustrates a perspective view of the battery case;

**FIG. 10** illustrates a side view in partial cross section of the medical diagnostic system;

**FIG. 11** illustrates a block diagram of the medical diagnostic system;

**FIGS. 12A, 12B AND 12C** illustrate an electrical circuit schematic diagram of the medical diagnostic system;

**FIG. 13** illustrates loading of the medical diagnostic system;

**FIG. 14A - 14I** illustrate component positioning during operation of the medical diagnostic system; and

**FIG. 15** illustrates display messages.

## 45 DESCRIPTION OF THE PREFERRED EMBODIMENTS

**FIG. 1** illustrates a perspective view of a portable pocketable glucose medical diagnostic system 10 including a disposable diagnostic reagent unit 12 as illustrated and later described in particular detail in **FIG. 4**. Externally visible components of the system 10 include front and back housing halves 16 and 18 respectively which enclose the electromechanical structure as later described in detail and, a battery case 20 and battery cover 22. An LCD or like visual readout 24 displays the glucose levels, time, battery

condition, stored values in memory, and other mode operational displays as later described in detail. Conductive rubber keypad buttons 26 and 28 position adjacent to the LCD readout 24. A actuator button 30 and a release button 32 locate on the top side of the glucose medical monitoring system 10 for subsequent cocking and releasing of a firing mechanism as later described in detail in the figures.

**FIG. 2** illustrates a perspective in view of the portable pocketable glucose medical diagnostic system 10, where all numerals correspond to those elements previously described. The flush rotatable protective dust cover 34 includes an interior mounted color reference calibration unit 36 and rotates on pivot blocks 38a-38b of Fig. 8, and conforms to the front housing 16 and the back housing 18 halves. The dust cover 34 includes a rectangular frame 39 for the accommodation of the interior mounted color reference color calibration unit 36 as also illustrated in Fig. 8. Later members 40 and 42 extend vertically from the interior side of the dust cover 34 and latch with the catches 44 and 46 at the upper edges of the rear and front housing halves 18 and 16 respectively. Vertical guide bars 48 and 50 at the ends of the rear and front housing halves 18 and 16 align along the outer surfaces of the latches 40 and 42 to align the dust cover 34 with the ends of the combined case halves 18 and 16. The rectangular frame 39 containing the color reference calibration unit 36 aligns in a rectangular hole 52 between vertical end bars 54 and 56 at the ends of the rear and front housing halves 18 and 16 respectively. An optics window 58 aligns with the rectangular hole 52 and also with the color reference calibration unit 36 when the dust cover 34 is rotated and latched in the closed position.

A clip 60 with a square mounting pad 62 frictionally engages a corresponding size hole 64 in the rear housing half 18.

**FIG. 3** illustrates a perspective view of the portable pocketable glucose medical diagnostic system 10, where all numerals correspond to those elements previously described illustrated in particular is the battery cover 22. The cover includes a back surface 22a and a side surface 22b between upper and lower curved surfaces 22c and 22d. Upper and lower curved surfaces 22c and 22d include end latch members 22e and 22f for snap engagement of the battery cover 22 within the battery case 20. The battery cover 22 members 22a, 22b, 22c and 22d form a carriage member into which a plurality of batteries 66a-66n are contained. The battery cover 22 and batteries 66a-66n engage within the battery case 20. The batteries 66a-66n are connected as later described in detail in Fig. 5.

**FIG. 4** illustrates a perspective view of the portable pocketable glucose medical diagnostic system 10 with a disposable diagnostic reagent unit 12 and a disposable lancet 14, where all numerals correspond

to those elements previously described. The disposable lancet 14 is inserted into the firing mechanism assembly 68 of Fig. 5 through an orifice 68. Orifice 68 aligns beneath the catches 44 and 46 and between the guide 4 bars 48 and 50 and in the end of the front and rear housings 16 and 18. After the disposable lancet 14 is inserted through the orifice 68, the disposable diagnostic reagent unit 12 is inserted between guide bars 48 and 50, in front of the orifice 68 and catches 44 and 46 and behind the vertical end bars 54 and 56. The disposable reagent pad 12 includes a hole 70 and reagent pad 72 and windows 74 and 76 for viewing of the blood soaked reagent pad 72 by internal electronic viewing as later described in detail. The hole 70 in the disposable diagnostic reagent unit aligns with the lancet needle and the windows 74 and 76 align with the optics window 58 for electronic viewing.

**FIG. 5** illustrates an exploded view in perspective of the portable pocketable glucose medical diagnostic system 10, where all numerals correspond to those elements previously described. The front housing half 16 includes a release cutout 78a a spring containment channel 80, a release button spring 82 between the spring containment channel 80 and the release button 32, a supported horizontally aligned firing mechanism track member 84, a spring seat 85 between the firing mechanism track member 84 and the top of the housing member 16, an optics head track member 86, side spacer bars 88 and 90, switch mounts 92a-92b and switch mount 94a-94b. A slotted cutout 96a and release cutout 78a align along the top inner edge of the front housing half 16 and a slotted cutout 96b and a release cutout 78b align along the top inner edge of rear housing half 18 to accommodate movement of the release button 32 vertically and the actuator button 30 horizontally. The rear housing half 18 also includes mirror like image elements of the firing mechanism track member 84 and the spring seat 85 and the optics head track member 86. The front housing half 16 also includes a piezo sounding device mount 98, a rectangular bracket 100, a bracket 102 for mounting of a conductive rubber keypad 104 containing buttons 26 and 28 and pivot hole 106 at the lower end. The rear housing half 18 also includes a pivot hole 108. Pivot holes 106 and 108 in housing halves 16 and 18 accommodate the pivot blocks 38b and 38a of the dust cover 34 respectively. A clear plastic display window 110 and the LCD panel 24 align and secure in the rectangular bracket 100 a foam pad 112 and an elastomeric LCD connector 114 align between the LCD panel 24 and electronics circuit board 116. The electronics circuit board 116 connects to an optics head 118, a switch 120, and a switch 122 through a flex cable 124. An optics spring retainer 126 and an optics spring retainer 128 align behind the optics head 118 as later illustrated. The optics head 118 aligns between the firing mechanism track member 84 and the

optics head track member 86 on the front housing half 16 and corresponding track members on the rear housing member 18. The firing mechanism assembly 68 aligns between the firing mechanism track member 84 and the top portion of the front and rear housing halves 16 and 18 and include the actuator button 30, a lancet carrier 130 a firing spring 132 and a return spring 134. A switch actuator 136 aligns between the optics head track member 86 and the bottom of the front housing half 16 and also between the corresponding members on the rear housing half 18. A positive and negative battery contact assembly 138 and 140 align and secure to the battery case 20. A user label 142 aligns in a label mount recess 144 on the rear housing half 18.

FIG. 6 illustrates the optics head 118 and associated components where all numerals correspond to those elements previously described. The assembly is illustrated on its side for clarity of illustration. An LED 146, an infrared LED 148 and a photo diode 150 secure within the optics head 118 with adhesive units 152 and are canted at an angle. The optics head includes ramped surfaces 150a-150b extending from the surface 152 of the cube like optics head 118. These ramped surfaces 150a-150b assist in the sliding action of the optics head 118 when the disposable diagnostic reagent unit 12 is inserted into the portable pocketable glucose medical diagnostic system 10. An optics cover 154 fits over and about the optics head 118. A thermister 156 attaches to the flex cable 124.

FIG. 7 illustrates an exploded view of the firing mechanism assembly 68 where all numerals correspond to those elements previously described. The lancet carrier 130 is the nucleus of this assembly and includes a cylindrical body 158, an interior lancet cavity 160, a longitudinal horizontally aligned slot 162 in the upper portion of the cylindrical body 158, a vertically aligned slot 162 across the cylindrical body 158 and intersections the longitudinal horizontally aligned slot 162, detented actuator bars 164 and 166 including rear detents 168a and 168b and forward detents 170a and 170b. The actuator button rests atop a hammer body 172. A hammer 174 aligns on one edge of the hammer body 172. A cam 176 for actuation of the switch 122 extends laterally to the side of the hammer body 172. The hammer body 172 and hammer 174 fit in and slide within the lancet cavity 160. A spring seat 178 in the form of a ring is molded about the circumference of the cylindrical body 158. The firing spring 132 fits over and about the cylindrical body between the spring seat 178 and the hammer body 172 as illustrated in FIG. 8. The return spring 134 seats between the spring seat 178 and the end of the housings 16 and 18 as illustrated in FIG. 8. A planar member 180 aligns between the actuator button 30 and the hammer body 172. At the continuous slot between the planar member 180 and the actuator button 30 rides

along slots 96a and 96b in the case halves 16 and 18 and serves to keep the hammer aligned in the lancet cavity 160.

FIG. 8 illustrates a perspective view of the dust 5 cover where all numerals correspond to those elements previously described. Illustrated in particular is the color reference calibration strip 36 which aligns to the rectangular frame 39. Pivot blocks 38a and 38b are integral to and extend inwardly from the pivot bar 10 members 184 and 186. Opposing pivot bar members 184 and 186 extend vertically from the main body 188 as to the latches 40 and 42. Pivot blocks 38a and 38b engage pivot holes 108 and 106 respectively of FIG. 5.

FIG. 9 illustrates a perspective view of the battery 15 case 20, where all numerals correspond to those elements previously described. Positive and negative battery contact assemblies 138 and 140 include 20 spring contactors 138a and 140a which frictionally engage a plastic securing plate 190 in the end of the battery case 20. The spring contactors 138a and 140a contact batteries 66a - 66n which are held in the battery cover 22 of FIG. 3. Wires 192 and 194 are electrically connected to and extend from the battery contact 25 assemblies 138 and 140 and connect to the electronics circuit board 116.

FIG. 10 illustrates a side view in partial cross section of the medical diagnostic system 10, where all numerals correspond to those elements previously 30 described. The cylindrical body 158 with the included and internally aligned hammer body 172 align between the top of the case, the firing mechanism track member 84, and the front position of the front and rear housing halves 16 and 18. Firing springs 132 and 35 return spring 134 both seat against opposing sides of the spring seat 178 on the cylindrical body 158. The return spring also seats around and about the material surrounding the orifice 68. The firing spring seats against the spring seat 85 of the hammer body 172, 40 slides within the lancet cavity 160, in the cylindrical body 158, and contacts engage and compresses the firing spring 132, when moved to the right in this illustration. The hammer body 172, including the cam 176 are actuated along longitudinal axis by the sliding of 45 the actuator button 30. As the cylindrical body 158 is positioned longitudinally the attached detented actuator bars 164 and 166, are longitudinally positioned to engage or disengage detents 168a - 168b and 170a - 170b, upon both sides of the planar member 180, on the release button 32. A spring 82, seats 50 in the spring containment channel 80 and against the interior of the release button 32 to spring the release button 32 outwardly. The release button 32 aligns in the release cutouts 78a and 78b also illustrated in FIG. 5. The actuator button 30, including the planar member 180 captures the edges of the case halves 16 and 18 adjacent to slotted cutouts 98a and 96b illustrated in FIG. 5. The cam 176 actuate the optics head 55

switch 122 at its rear most travel. The optics head 118 aligns along the side spacer bars 88 and 90 and the optics head track member 86 of the front housing half 16 and corresponding members on the rear housing half 18. A spring alignment post 196 is included on the rear side of the optics head and another spring alignment post 198 is included on the optics spring retainer 126. A spring 128, aligns over the spring alignment posts 196 and 198 to slideably retain the optics head 118, in a position to the right of the spring and against the vertical end bars 56 and 54 found also in Fig. 5. A switch actuator bar 136 is actuated against the optics head switch 120 as later described in detail.

FIG. 11 illustrates and electrical block diagram 200 of the medical diagnostic system, where all numeral correspond to those elements previously described and as now described in Figs. 12a, 12b and 12c.

FIGS. 12A, 12B and 12C illustrate the electrical circuit schematic package diagram 201, including the digital display 202, clock and alarm switches 203 and 204, light emitting diodes 205 and 206, photo diode 207, strip switch 208, lance switch 209, piezo electric beeper 210, and batteries 211 and 212. A high gain op-amp 213 including, op-amp feedback capacitor 214, and op-amp pull-up resistor 215 are for the amplifier circuit. A to D converter 216, inverter transistor 217 for clock for A to D converter 216, pull-up resistor 218 for inverter 217, inverter transistor base drive resistor 219, voltage reference regulator integrated circuit 220, voltage reference regulator input bypass capacitor 224, output adjustment resistor 222, output adjustment potentiometer 223, output filter capacitor 221, voltage reference resistor 225, and voltage reference diode 226 are for the A to D conversion of the several colorimetric change of the reagent and the voltage reference regulator. Switching transistor 237 for LED 205, switching transistor 238 for LED 206, switching transistor base drive resistor 273, and switching transistor base drive resistor 274, are for switching the LED's, LED 205 brightness adjustment resistor 240, LED 206 brightness adjustment resistor 239, are for compensating the LED's. RC oscillator circuit capacitor 245, RC oscillator circuit resistor 246, reset capacitor 243 and reset resistor 244 are for the microprocessor 249. Piezo electric beeper impedance load resistor 232, microprocessor pull-down resistor 241, microprocessor pull-down resistor 242, microprocessor pull-down resistor 247, microprocessor pull-down resistor 248, analog power switching transistor 271, switching transistor base drive resistor 270, reverse voltage protector diode 272, 32.768 KHZ crystal for timer 255, timer current limiter resistor 254, crystal oscillator capacitor 256, and crystal oscillator capacitor 257 are for the microprocessor 249. A to D converter 233 for temperature variations, thermistor 273, and voltage divider resistor 234 are for the temperature sensing circuit. LCD bias resistors 261-264

and LCD bias capacitors 258-260, are for LCD display 202. Serial data output enable jack 275, serial data output jack 281, and serial data clock jack 282 are for external connections such as to a personal computer.

5 Bypass capacitors 205-253, low battery and very low battery comparators 230, and comparator voltage dividers 227-229 are for power supply circuitry. EEPROM power transistor 266, and EEPROM power base resistor 265, control power to serial EEPROM 267 and 268.

10 The operation of the electrical circuitry of FIGS. 12a and 12b is now described in detail. LED 205 is the light source that illuminates the reagent chemistry area. The reagent chemistry changes color in proportion to the amount of glucose in the blood. The light from LED 205 reflects off the chemistry and is sensed by photodiode 207. This signal is amplified by a high gain op-amp 213, and then sent to the input of the analog to digital converter 216. The analog signal is converted to a digital signal for use by microprocessor 249. The software algorithms in microprocessor 249 processes this information, and then outputs a blood glucose measurement to the liquid crystal display 202.

15 25 LED 206 is the light source that illuminates the lot to lot indicator on the medical diagnostic system 10. This provides information to the microprocessor 249 to correct for variations in different lots of chemistry. The lot to lot indicator also is used to determine if 30 blood has completely covered the reagent chemistry. The reflected light from LED 206 is sensed by photodiode 207, and the signal is sent to the microprocessor 249 in the same way as light reflected from LED 205.

35 40 Voltage reference regulator 220, provides a reference voltage for the medical monitoring system circuitry. The reference voltage is used by the analog to digital converter 216, the low battery detection comparators 230, temperature A to D converter 233 and also to keep the LED outputs constant.

45 50 Comparators 230 are used to provide a low battery and very low battery signal to the microprocessor 249. Switching transistor 271 is used to control the power to the analog circuitry which is turned on only when the photodiode 207 sensing circuits are active. A crystal 255 provides a precision clock to the microprocessor 249 for the various timing functions. Switch 209 is used to initiate a blood glucose measurement sequence by the medical monitoring system 10. Switch 208 provides the microprocessor 249 with a signal to tell when a reagent unit 12 is inserted. 55 Switches 203 and 204 are used to set the clock and four alarms on the medical monitoring system 10. The piezo electric beeper 231 provides an audible beep to indicate test progress or error conditions. Thermistor 273 with A to D converter 233 provide temperature correction input data to microprocessor 249 to correct for ambient temperature variations which may occur

in the user's environment. EEPROMs 267 and 268, provide non-volatile memory storage for alarms, saved glucose reading and various coefficients used in microprocessor 249, calculations.

**FIG. 13** illustrates the loading of the medical diagnostic system 10, mode of operation with a reagent unit 12, where all numerals correspond to those elements previously described.

**FIG. 14a - 14i** illustrate the component positioning and mode of electromechanical operation for the medical diagnostic system 10, where all numerals correspond to those elements previously described.

**FIG. 14a** illustrates normal position of the elements;

**FIG. 14b** illustrates the actuator button pushed into loading position;

**FIG. 14c** illustrates the button released and locked in a loading position;

**FIG. 14d** illustrates a lancet inserted while in a loading position;

**FIG. 14e** illustrates a button pressed into a cocked position;

**FIG. 14f** illustrates a strip inserted;

**FIG. 14g** illustrates a full extended position during penetration;

**FIG. 14h** illustrates a returned to normal position after penetration; and,

**FIG. 14i** illustrates the button pressed into an ejection position for removing and ejecting the lancet.

In operation, push the sliding button forward. Insert the lancet into the carrier tube. Remove the cap, press the release button. With the optic cover closed, pull back the sliding button. Open the optic cover and insert the reagent unit into position. Press a finger firmly on the reagent unit collector. Press the release button and squeeze the finger for hanging a drop of blood. Place on wick in the blood bowl. Read the glucose value displayed after 90 seconds, and record the results. Pull out and discard the used reagent unit. Push the sliding button forward to eject the used lancet, and close the optic cover.

**FIG. 15** illustrates display messages generated by algorithms in the microprocessor for display in the LCD.

The term "chemistry" used herein refers to a chemical reagent or to an indicator or signal means which operates in a chemical manner.

One purpose of the lot-to-lot identification window, where present, is for differences in the manufacturing of material and calibrates the system prior to reading the reagent.

Although the present invention is described above with particular reference to the use of color to represent the quality, quantity or condition being detected and/or measured, the invention includes other forms of signal, for example visual or audible.

Optional features of the invention which are of

particular interest are as follows:

System:

- 5 – the blood reagent chemistry is glucose oxidase chemistry
- includes means to return said hammer means to an original rest position
- 10 – includes cocking means connected to said hammer means
- includes means providing a light-tight enclosure of said housing of said unit
- includes piezoelectric audio means connected to said microprocessor means for beeping on predetermined conditions
- 15 – said microprocessor means includes means for displaying operational messages on said display means
- said microprocessor means includes means for displaying a plurality of previous readings
- 20 – said microprocessor means includes means for self-calibration
- includes a dust cover adjacent to said optical measurement means, and said dust cover including a calibration chart means for reading by said optical measurement means
- said housing is rectangular
- includes means for clipping into a shirt pocket of a user
- 25 – includes means in said housing for battery replacement
- includes switch means on said housing and connected to said microprocessor means for setting and displaying time
- includes means in said microprocessor to display numerical values in the English system and metric system
- 30 – includes a dust cover means rotatable on said housing about said optical measurement means
- includes calibration means internally mounted on said dust cover means for reading by said optical measurement means
- 35 – said microprocessor is 4 bit
- includes means in said microprocessor for temperature compensation
- 40 – includes means in said microprocessor for detecting an adequate fluid sample to said blood reagent chemistry
- said reagent chemistry is a solid
- said optic measurement means comprises two LED's about a photodiode
- 45 – said housing is of a size to be comfortably accommodated by a user's hand
- 50 **Process**
- said analysis is for blood glucose
- 55 – comprises reading the back side of said reagent chemistry
- comprises filtering at least one blood compo-

ment before transporting blood to said blood reagent chemistry.

### Claims

1. A system for extraction and analysis of a component of a liquid, said system receiving a disposable diagnostic reagent unit which exhibits a color change on sensing a predetermined component in the liquid, said system comprising:
  - a. optical measurement means including a light source and light sensor for measuring light emanating from said source and reflected by reagent chemistry in said unit and having an optical characteristic proportional to the component of the liquid to be measured after transporting when liquid to said reagent;
  - b. said optical measurement means generating an electrical signal responsive to a change of said reagent chemistry and therefore also to the component to be measured;
  - c. microprocessor means processing said generated electrical signal;
  - d. display means responsive to said processed signal providing a visual readout representative of the analysis on said display means in said housing member; and
  - e. means for removably receiving a disposable diagnostic reagent unit including said reagent chemistry.
2. A system according to Claim 1, wherein the liquid is blood and said unit exhibits a color change on sensing a pre-determined condition and wherein the system has a housing member including a spring actuated hammer means in said housing member and a disposable lancet removably positioned on said hammer means.
3. A system according to Claim 1 or 2, which includes said disposable diagnostic reagent unit.
4. A system for operative connection to a test device which device tests a liquid for the presence therein of an analyte, the system comprising:
  - an optical measurement means including a light source and a light sensor for measuring light emanating from said source and light received from the test device and having an optical characteristic related to the presence of, or the quantity of, the analyte;
  - generating means to generate an electrical signal responsive to said optical characteristic;
  - microprocessor means to process said generated electrical signal;
  - display means responsive to said process-

sed signal and providing a visual display; and  
means for removably receiving the test device.

5. A system according to Claim 4, which includes said test device.
6. Hand-held pocketable medical monitoring diagnostic system for extraction and analysis of a component of blood in a body, said system comprising:
  - a. pocketable housing member including a spring actuated hammer means in said housing member and a disposable lancet removably positioned on said hammer means;
  - b. optical measurement means including a light source and light sensor for measuring light emanating from said source and reflected by blood reagent chemistry having a color optical characteristic proportional to the component of the liquid to be measured when in contact with the liquid;
  - c. said optical measurement means generating an electrical signal responsive to a color change of said blood reagent chemistry and therefore also to the component to be measured;
  - d. microprocessor means processing said generated electrical signal;
  - e. display means responsive to said electrical signal to provide a visual readout representative of the analysis on said display means in said housing member;
  - f. means for removably receiving a disposable diagnostic reagent unit; and,
  - g. disposable diagnostic reagent unit for operative engagement in said receiving means of said glucose medical monitoring diagnostic system, said unit including housing, means for operatively connecting said unit to said receiving means of said system, said blood reagent chemistry supported within said housing, at least one opening in said housing for providing an opening for a puncturing means to pass through and providing for a liquid substance to be transported to said blood reagent chemistry, means for transporting said liquid substance to said blood reagent chemistry, and aperture means through said housing for reading a portion of said blood reagent chemistry whereby said blood reagent chemistry yields a responsive optical characteristic which is read by said optical measurement means, processed by said microprocessor means, and displayed by said display means thereby yielding a numerical value of a diagnostic condition.

7. A system according to Claim 6, wherein said opening provides for transfer of a liquid substance to said reagent means.

8. A system according to Claim 6 or 7, which includes calibration means supported in an aperture next to said reagent means in said housing. 5

9. A system according to Claim 6, 7 or 8, wherein said transporting means is a wicking material. 10

10. A system according to any of Claims 6 to 9, which includes a calibration sensor for sending visual calibration color at predetermined times as sequenced by said microprocessor means. 15

11. A system according to Claim 6, which includes cocking means connected to said hammer means, and said microprocessor means includes means for processing a plurality of steps on cocking said hammer means. 20

12. A system according to Claim 6, which includes means in said microprocessor to verify lot-to-lot (LOT/LOT) reagent specific calibration codes of said calibration means. 25

13. A process for measuring a component quality comprising the steps of:  
 a. sliding a disposable diagnostic unit into a hand-held pocketable medical system; 30  
 b. engaging the medical system including said unit against an individual's skin;  
 c. puncturing the skin with a needle spring-biased in said unit; 35  
 d. transporting blood from outside the individual's skin for qualitative analysis by blood reagent chemistry carried by said unit; and  
 e. reading the qualitative results on a display of said system. 40

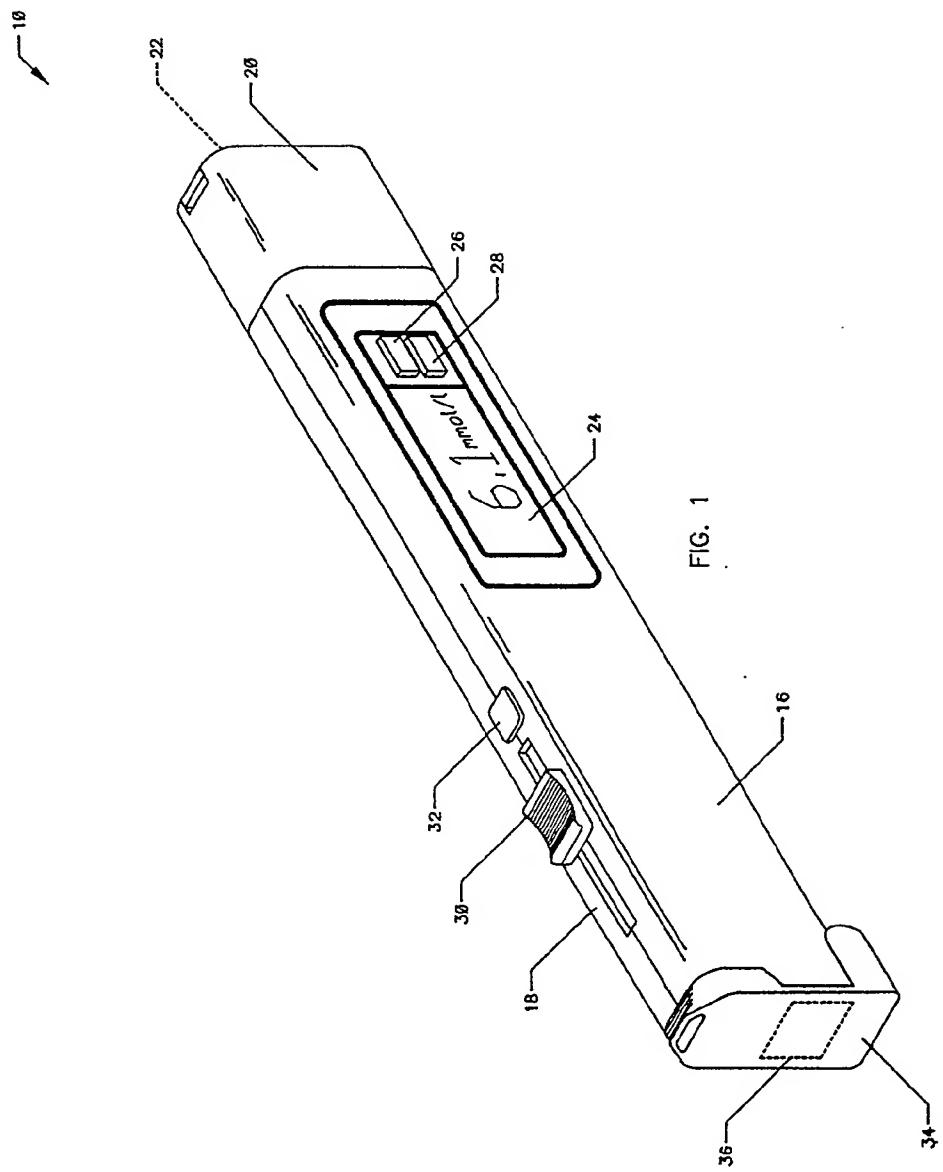
14. A process according to Claim 13, which includes calibrating said system prior to sensing a qualitative result. 45

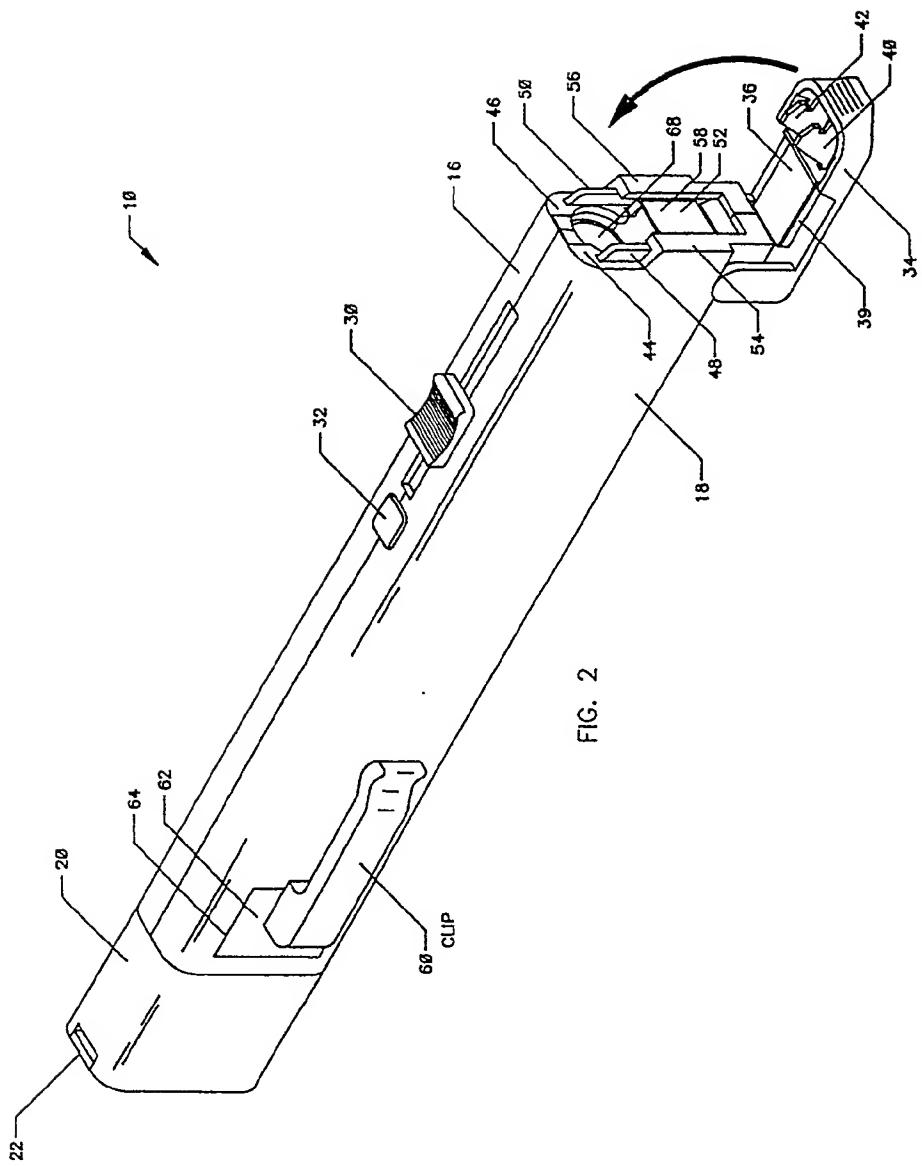
15. A process according to Claim 13, which includes the step of reading the previous qualitative results.

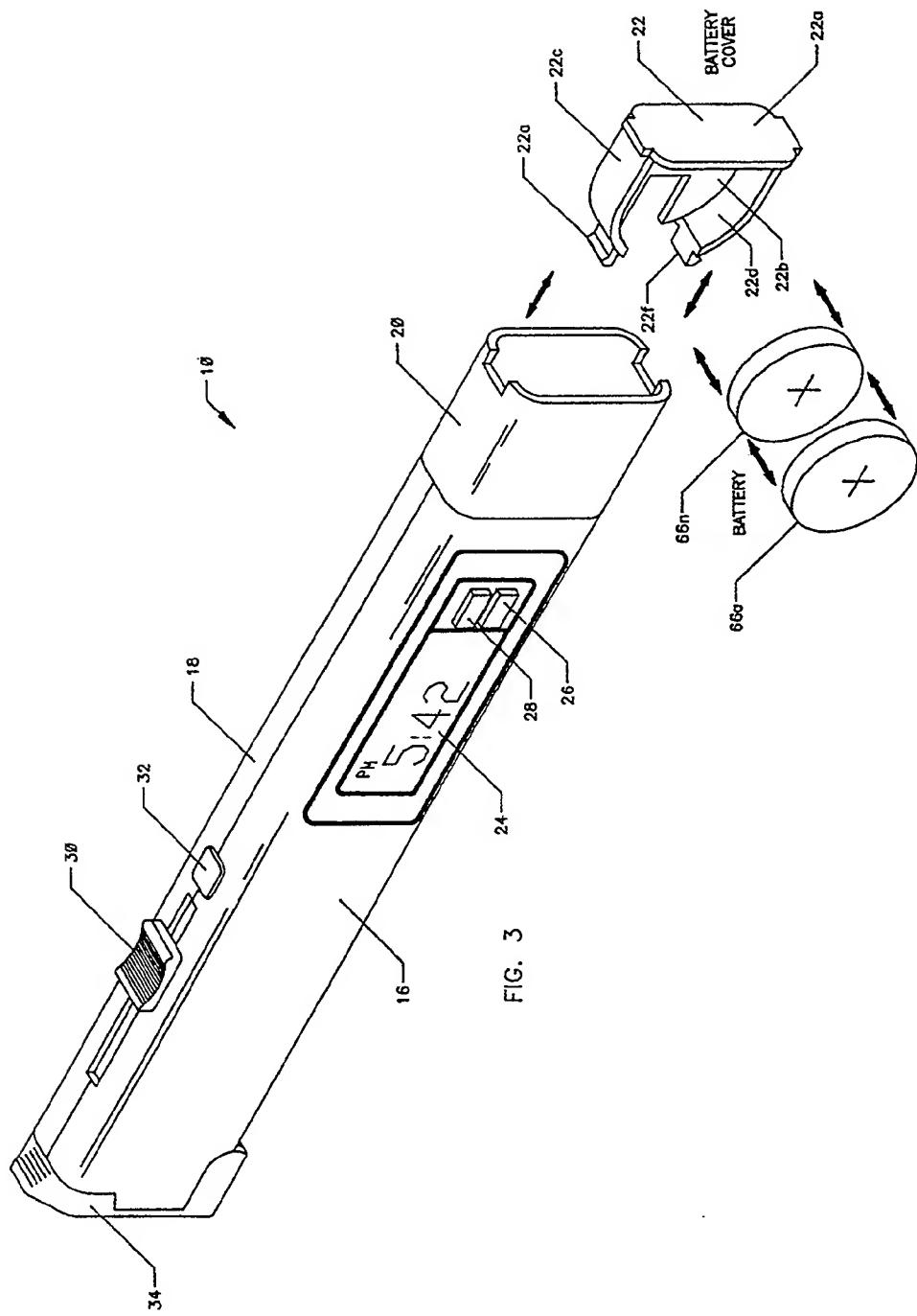
50

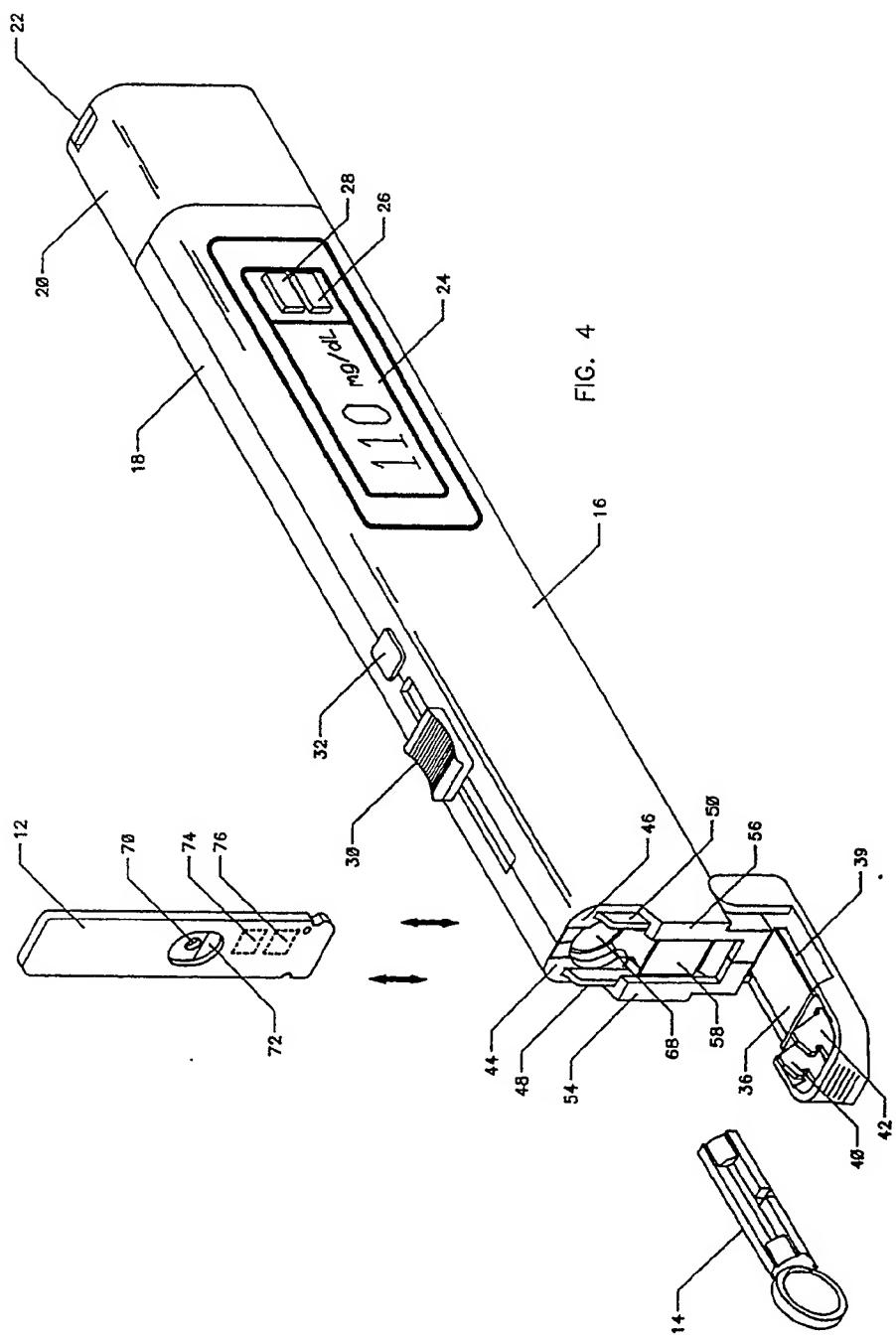
55

11









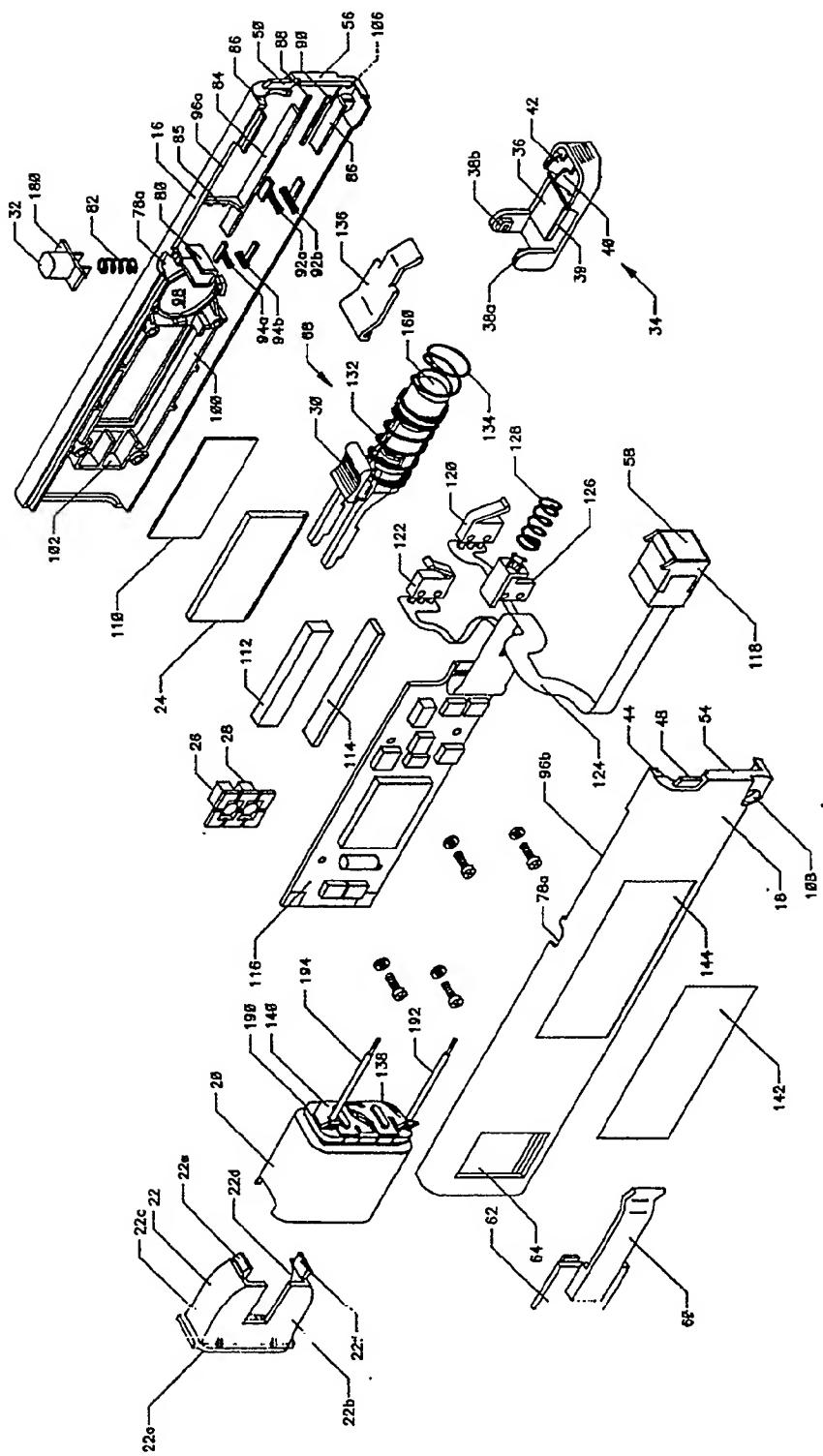


FIG. 5

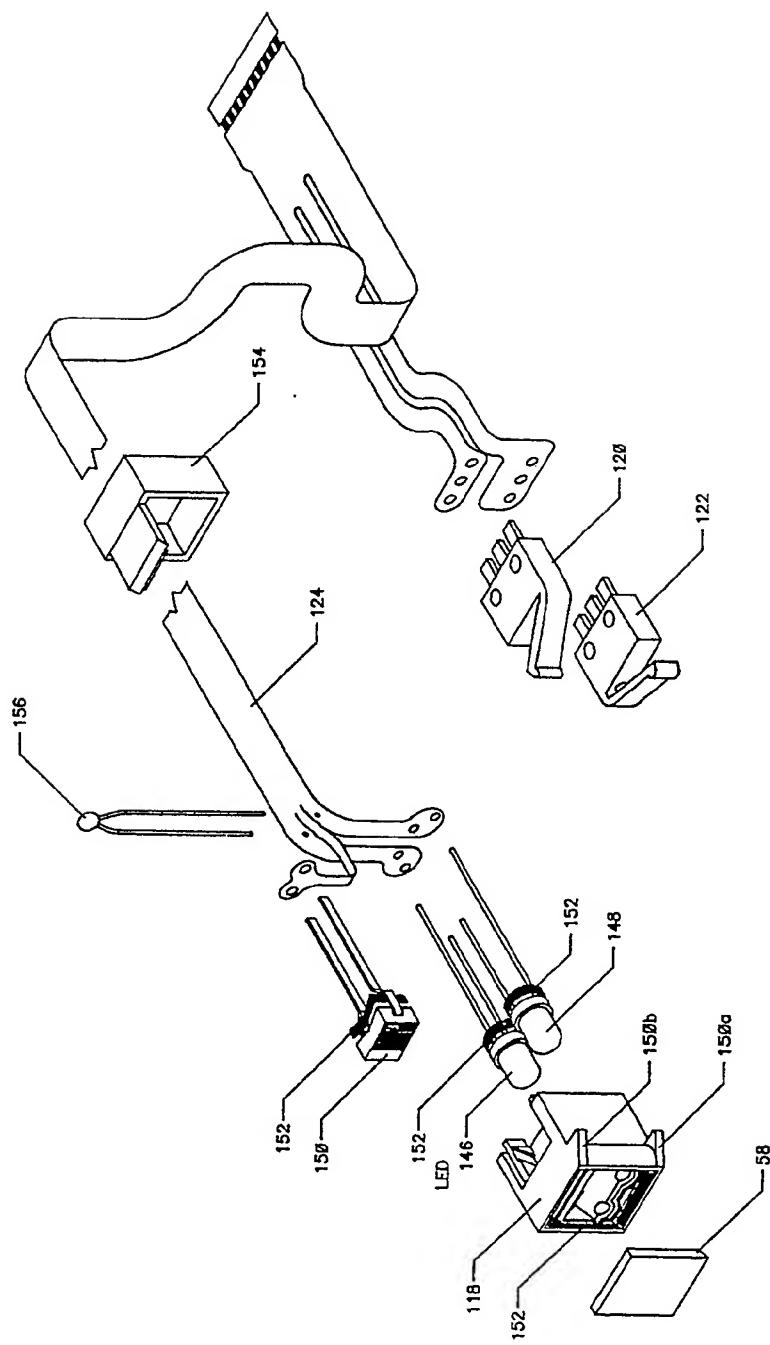


FIG. 6

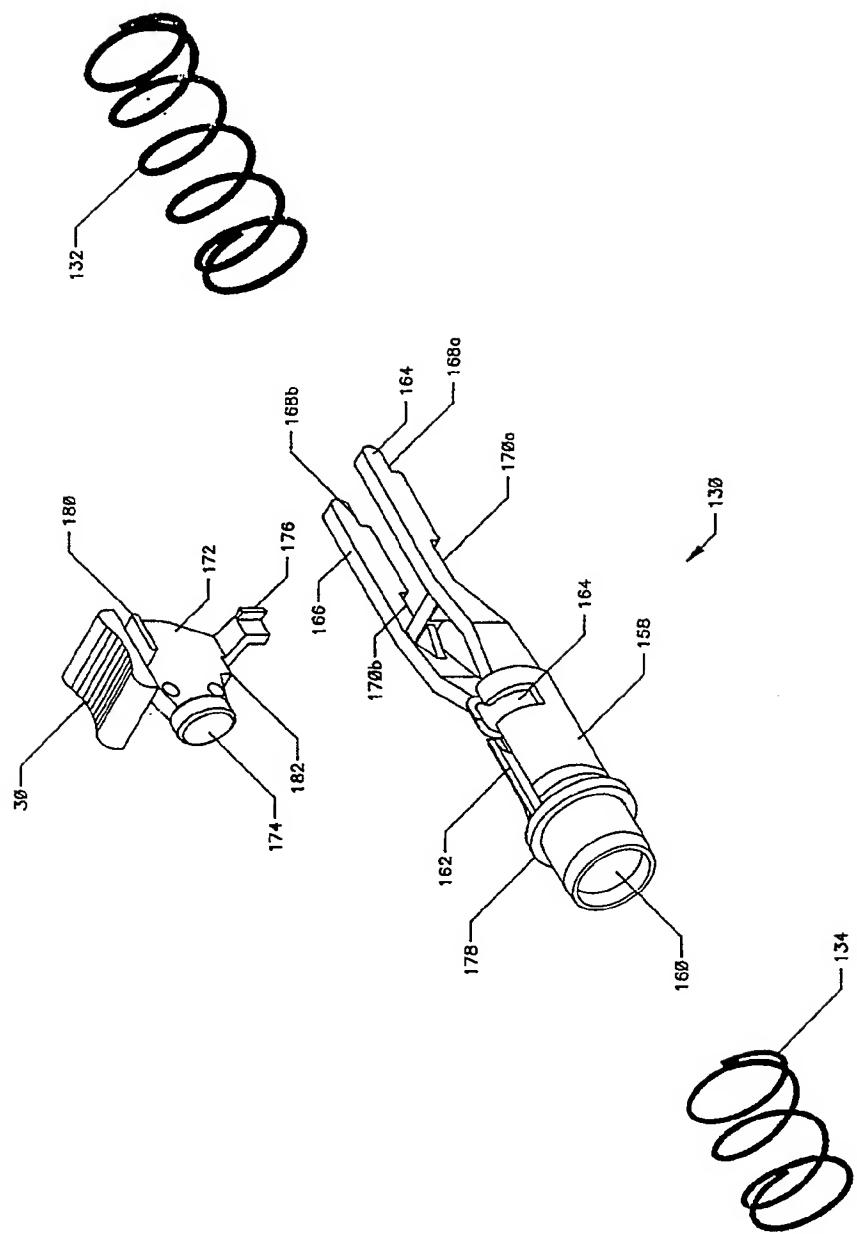


FIG. 7

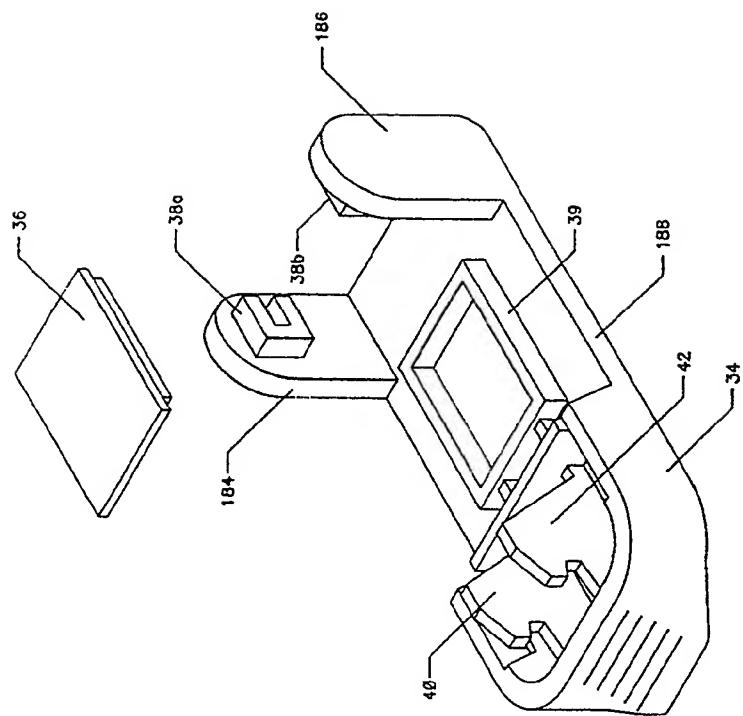


FIG. 8

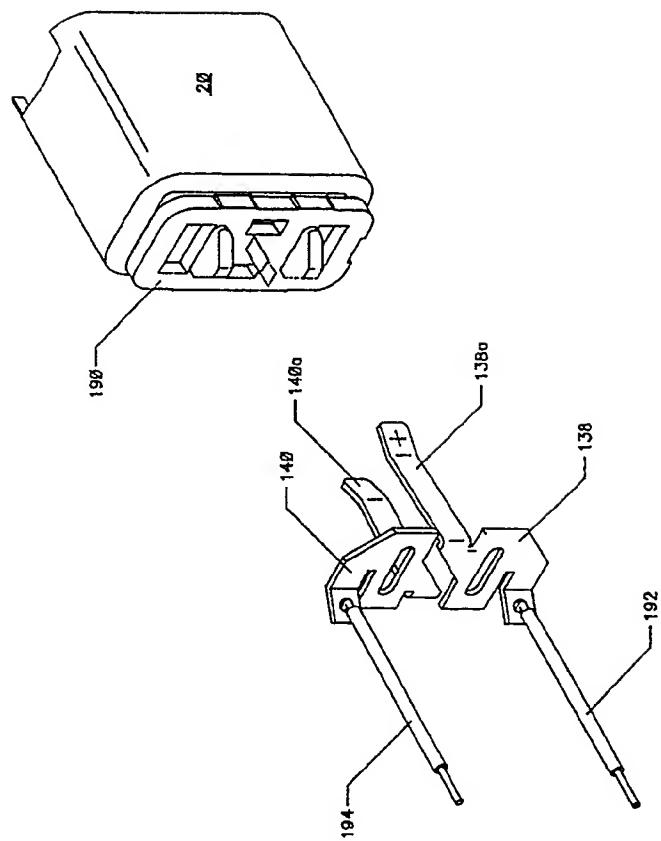


FIG. 9

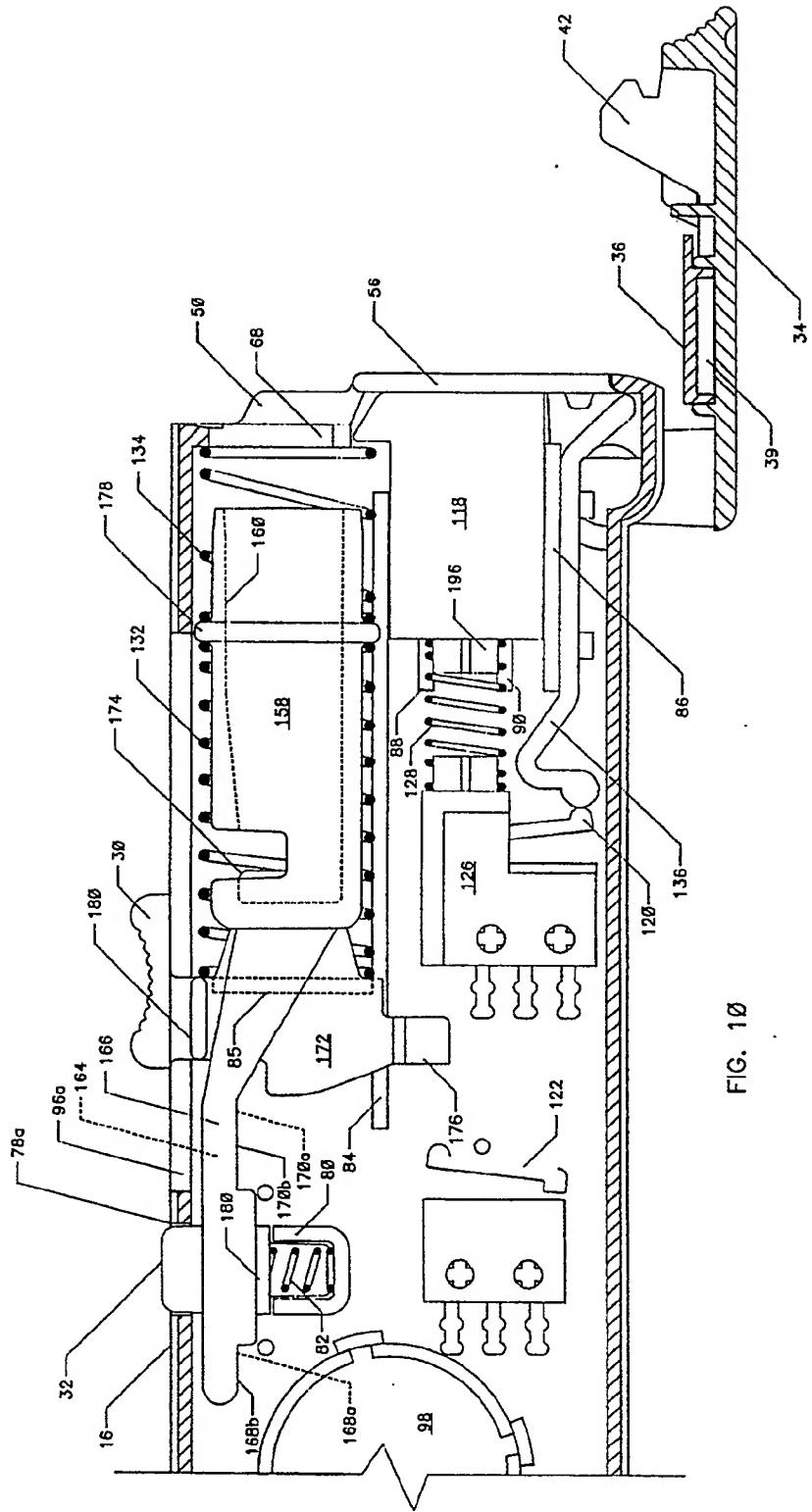
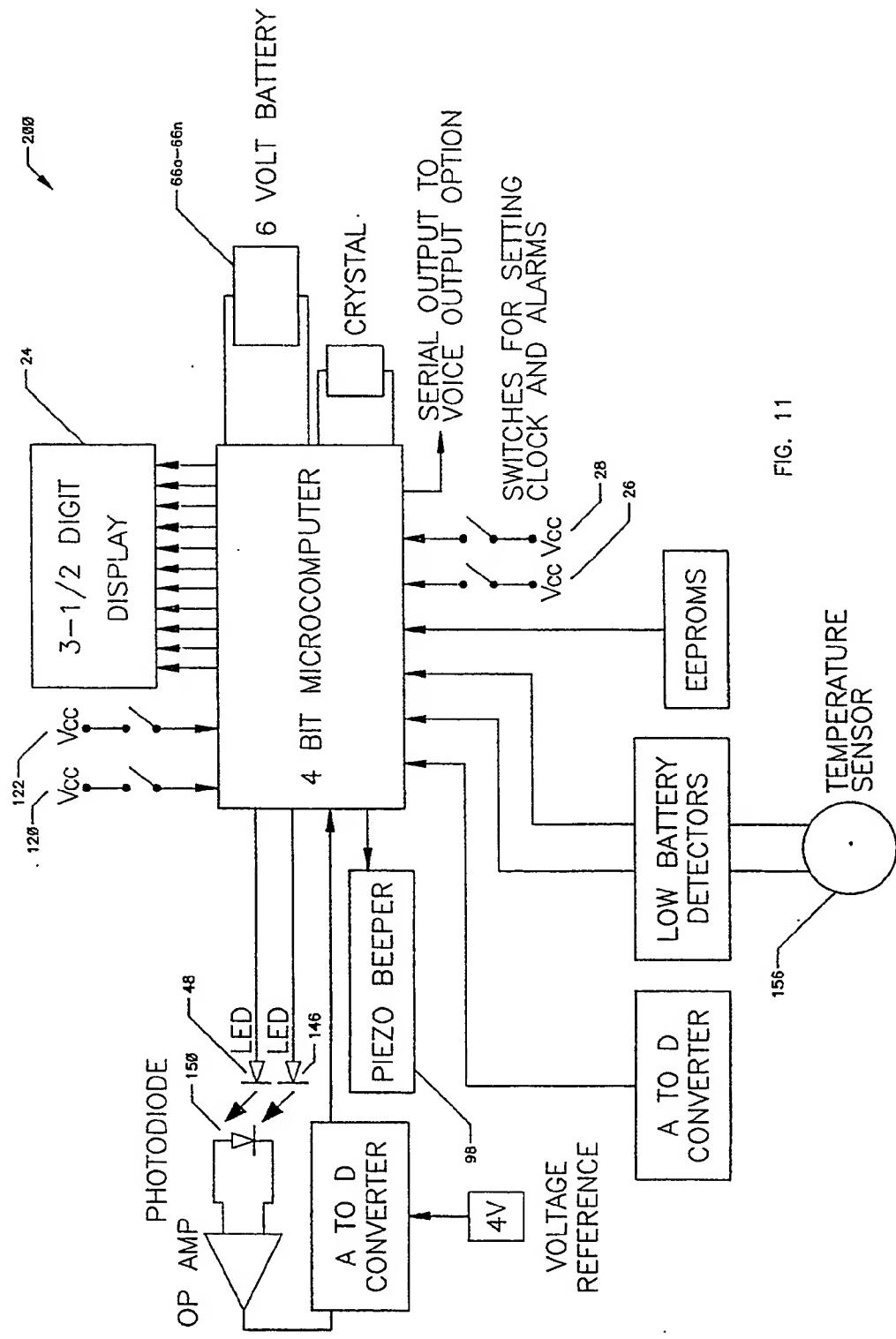


FIG. 10



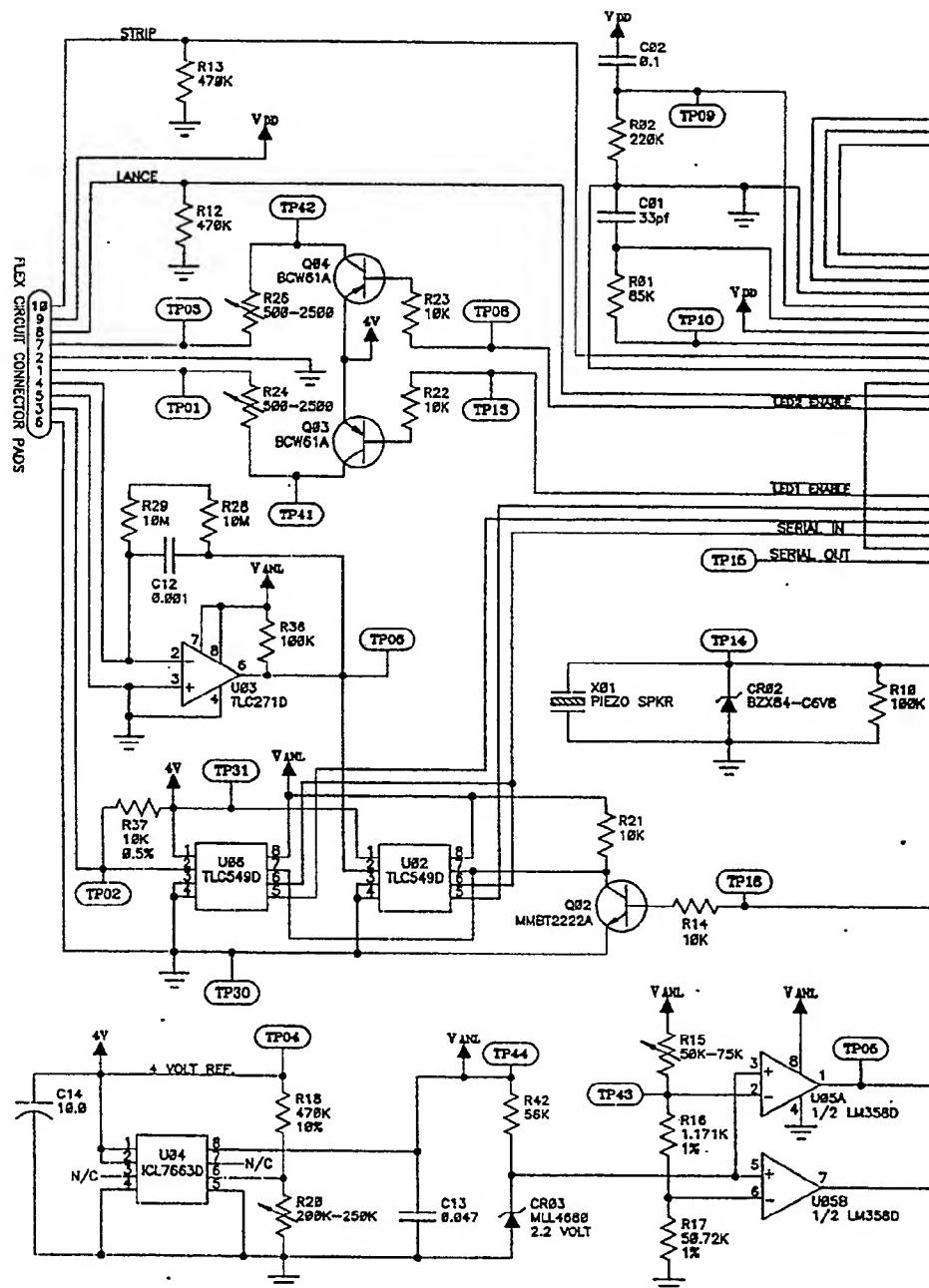
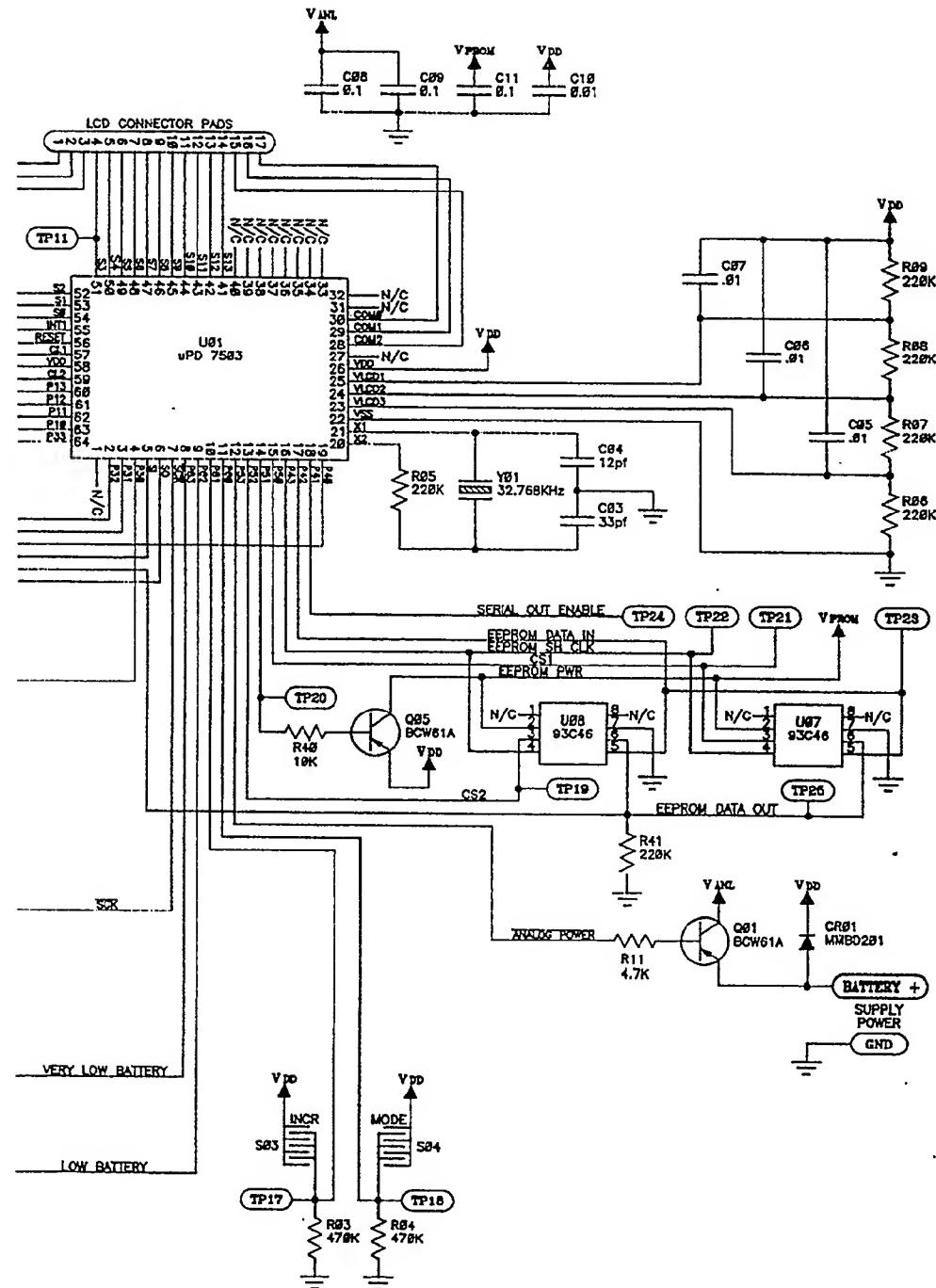


FIG. 12a



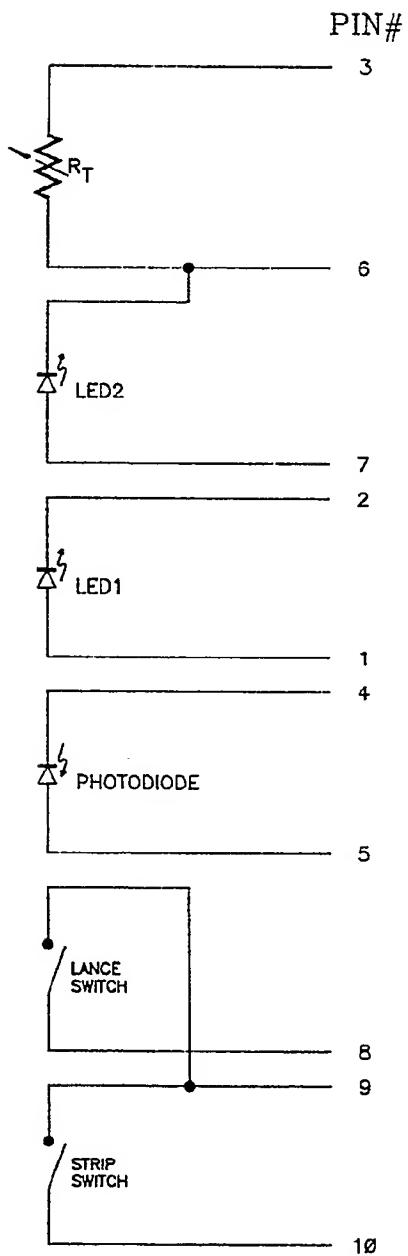
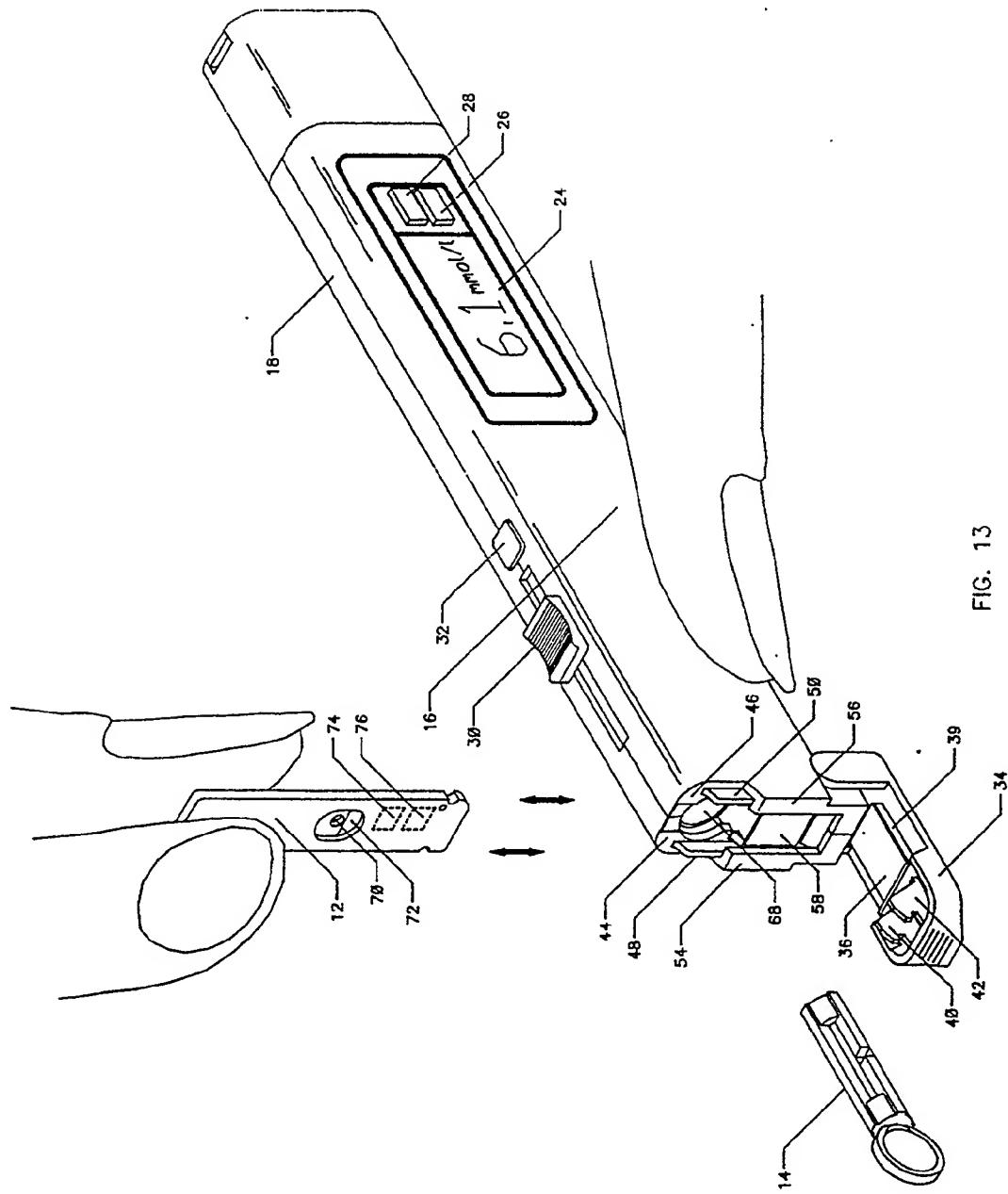


FIG. 12c



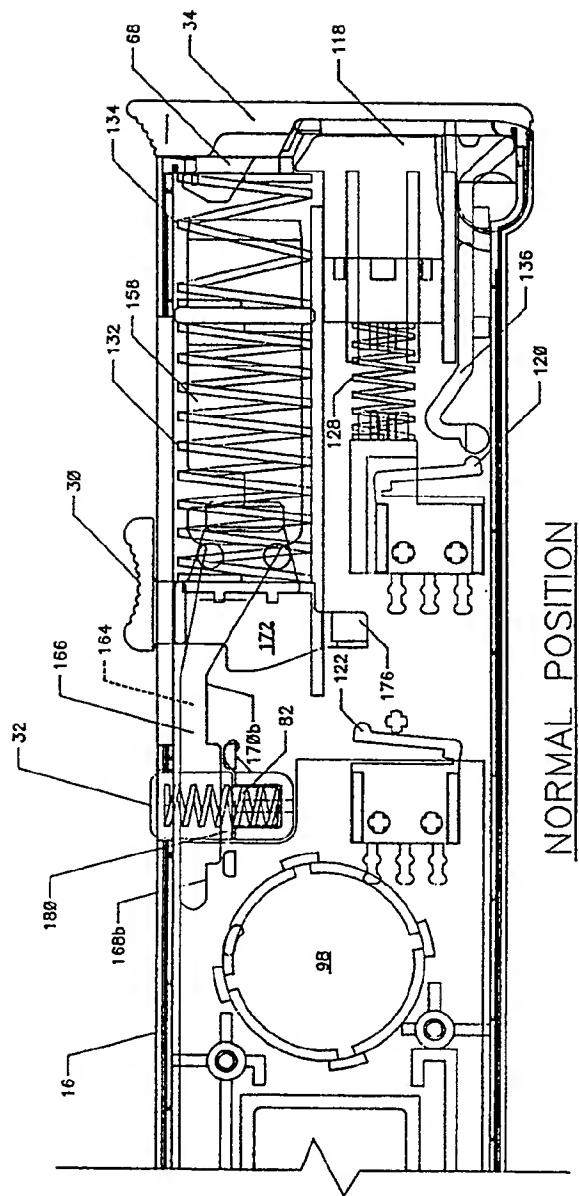


FIG. 14a

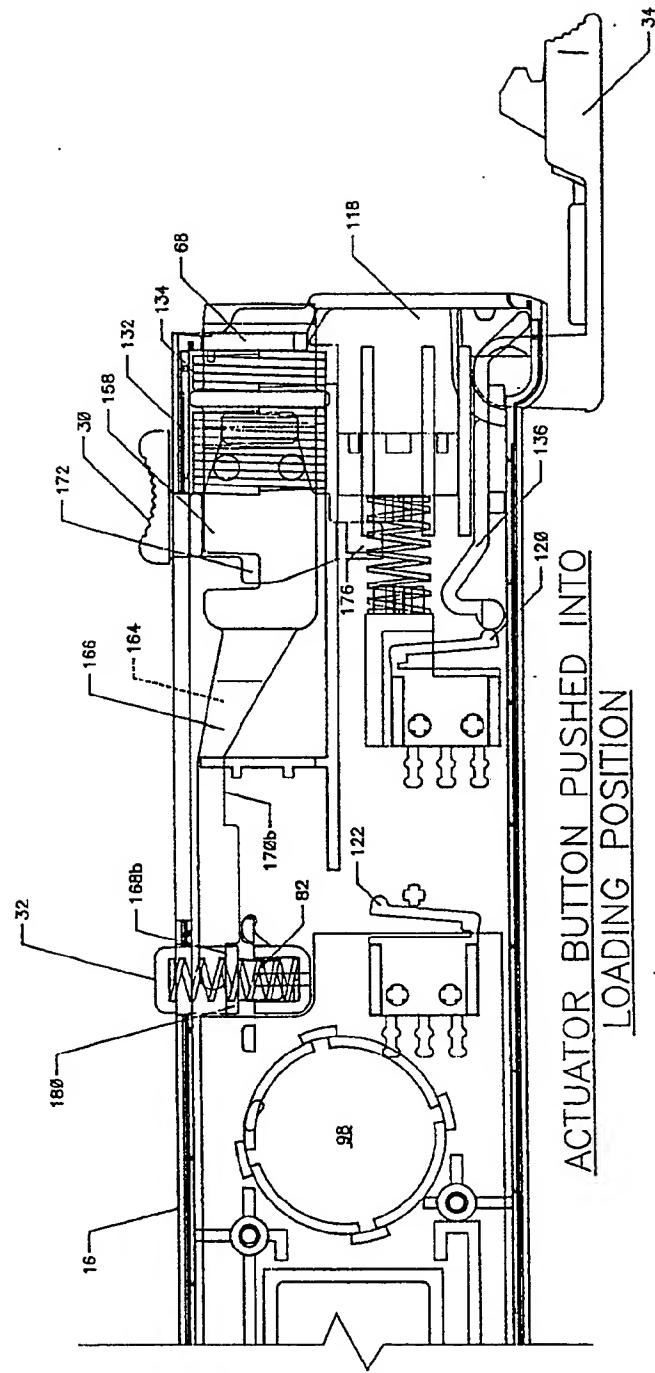


FIG. 14b

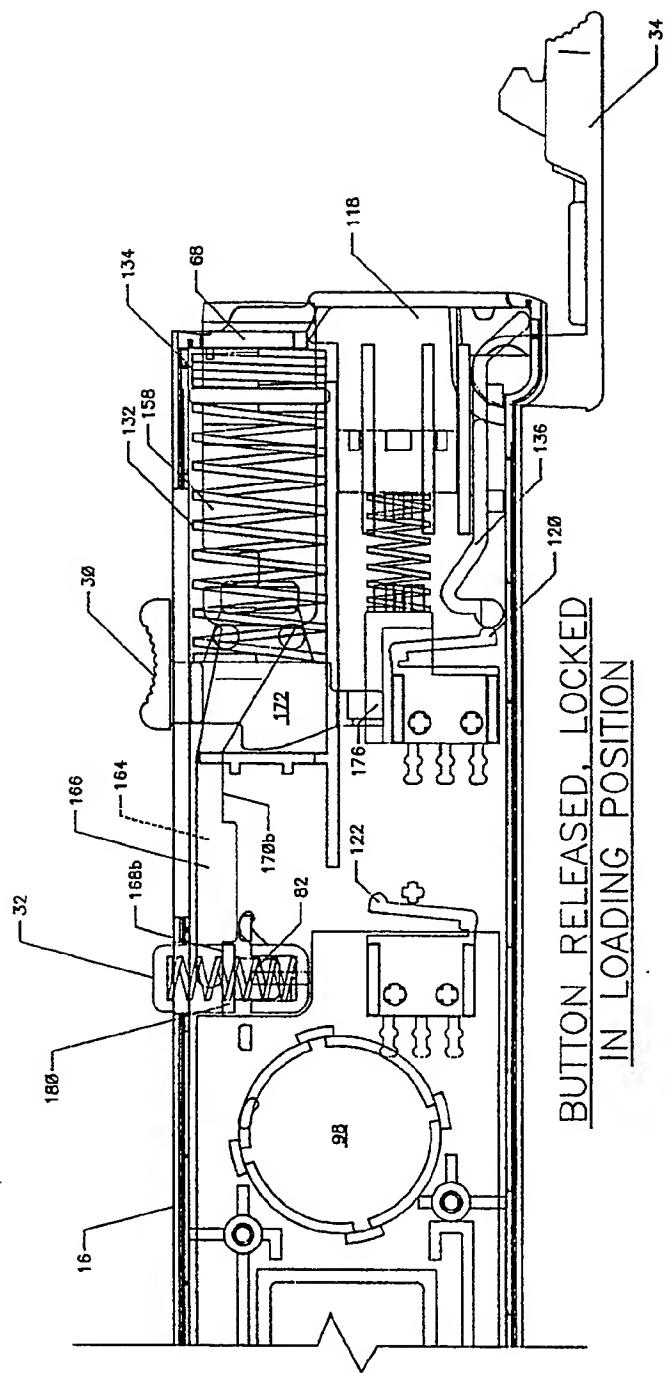


FIG. 14c

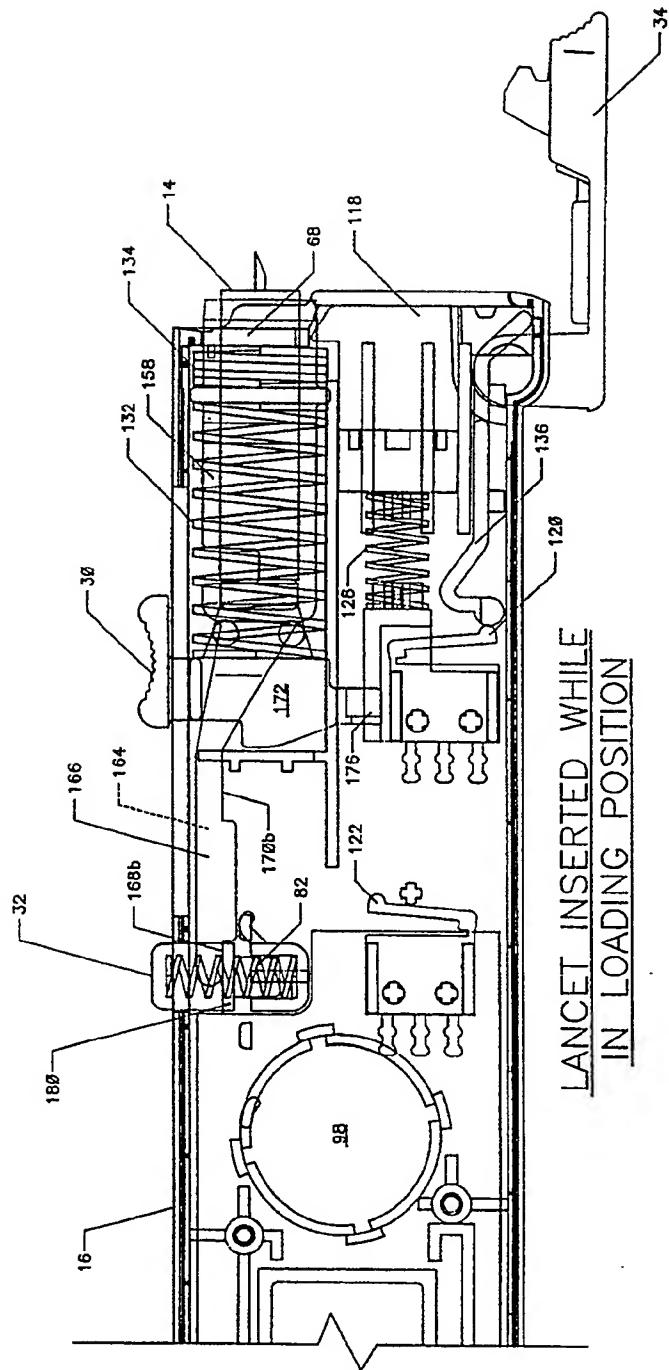


FIG. 14d

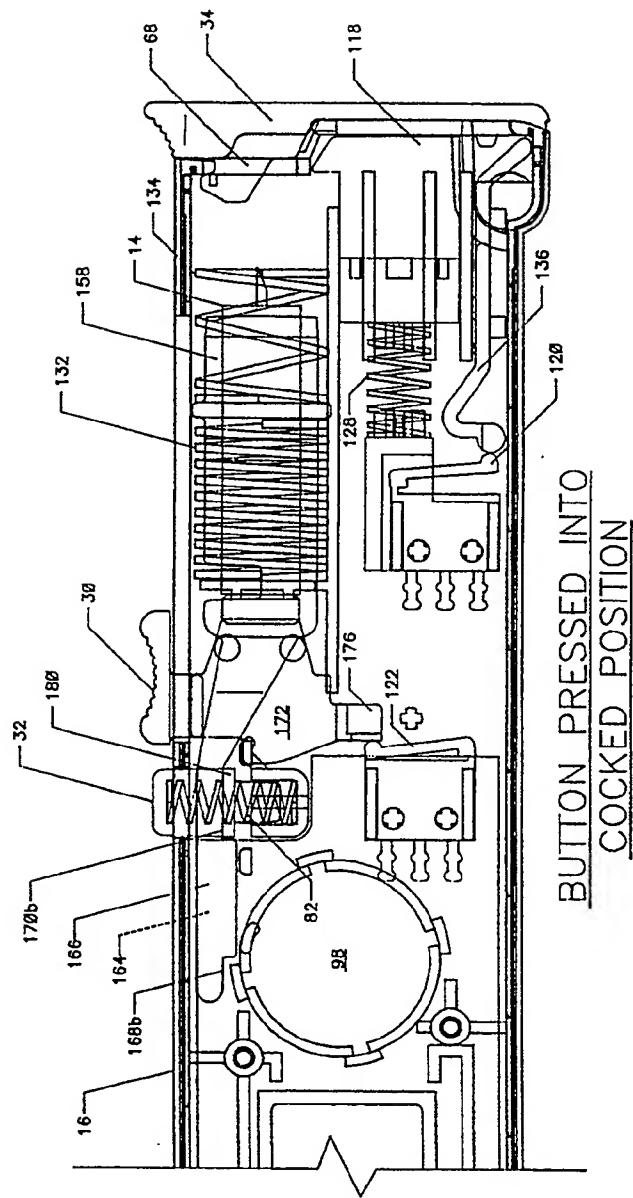


FIG. 14e

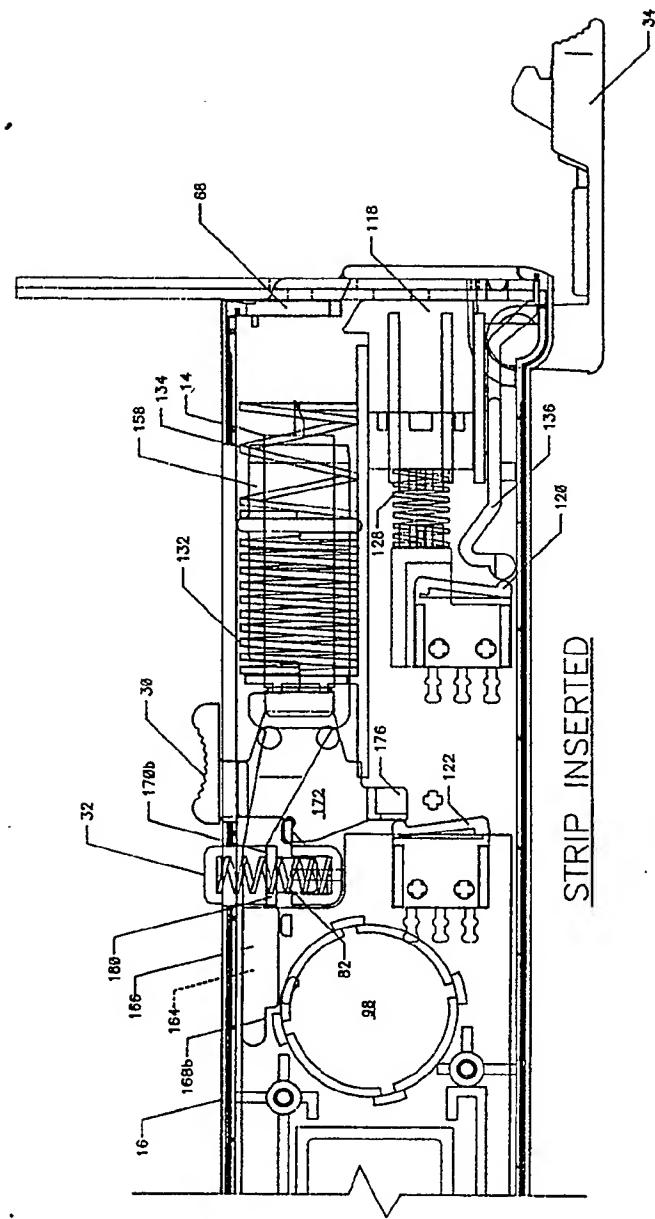


FIG. 14f

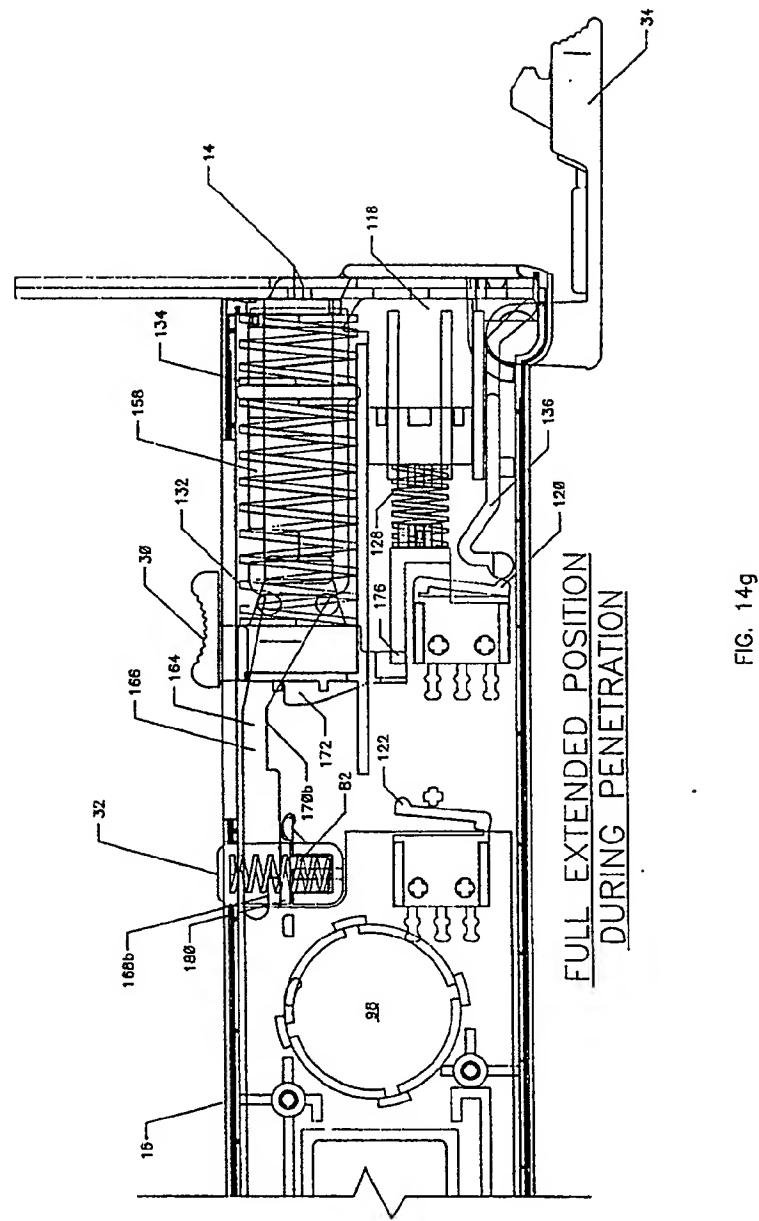


FIG. 14g

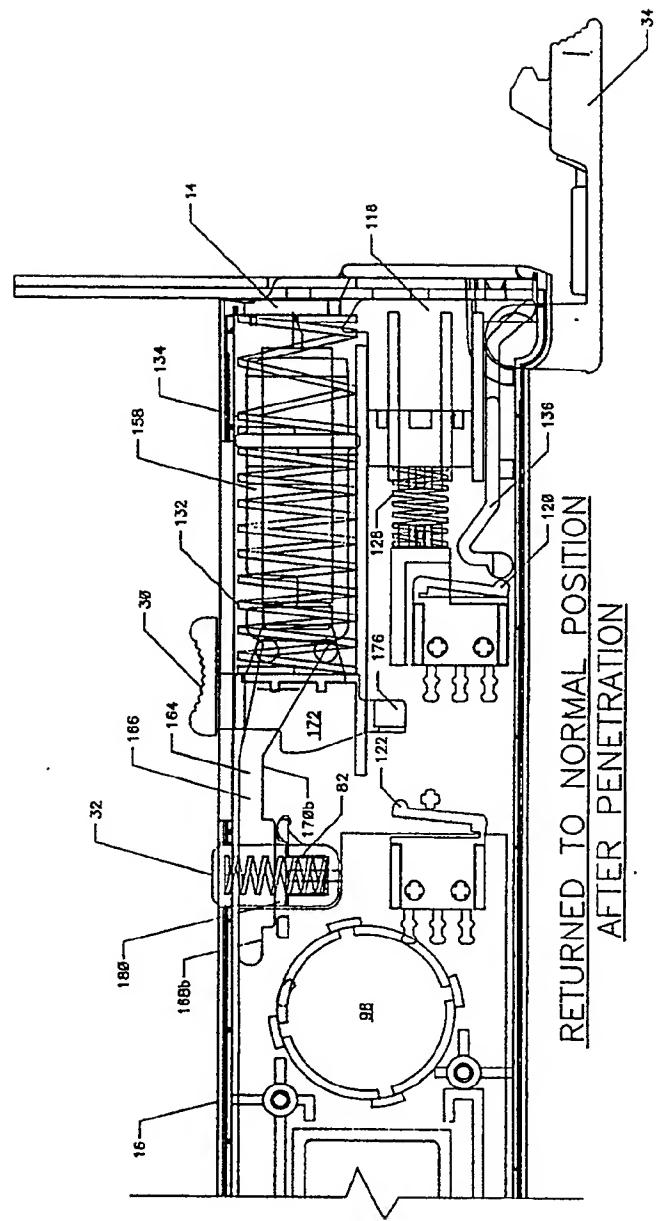


FIG. 14h

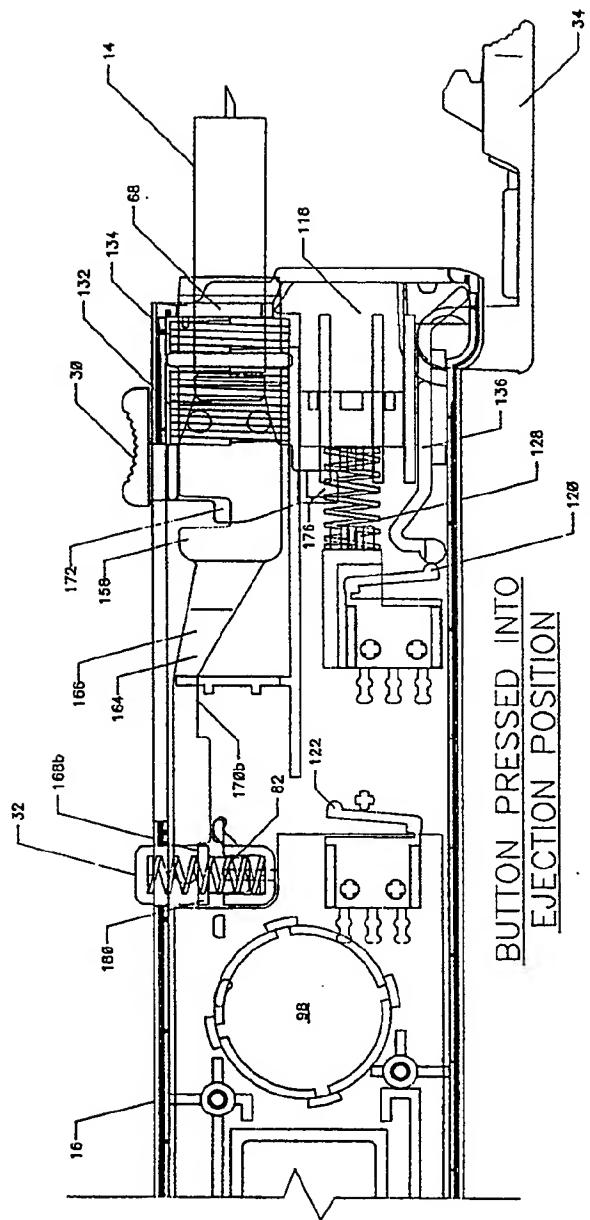


FIG. 14i

## DISPLAY MESSAGES

**H0E**

Temperature too high. Begin again at 63-95°F.

**---**

Batteries are low, meter will function properly. Replace Batteries soon.

**E-1**

(Continued)

- Dirt on Optic Window or Calibration Pad. Clean, begin again.
- Electronic malfunction. See Instruction Manual.

**E-3**

(Continued)

- ANSWER strip inserted upside down. Begin again with same ANSWER strip.

**E-6**

Too little blood applied to complete countdown. Repeat test with new ANSWER strip.

**CLd**

Temperature too low. Begin again at 63-95°F.

**EEE**

Batteries too low. Replace immediately.

**E-4**

- Too little blood applied within the time allotted to initiate countdown. Begin again with new ANSWER strip.
- No blood applied. Begin again with same ANSWER strip.

**LLL**

Result lower than 40 mg/dL. See Instruction Manual.

**E-0**

Electronics error. See Instruction Manual.

**E-2**

ANSWER strip dislodged or removed after firing Lancet. Begin with same ANSWER strip.

**E-7**

ANSWER strip not inserted within 3 minutes after Sliding Button was pulled back. Begin again with same ANSWER strip.

**HHH**

Result higher than 400 mg/dL. See Instruction Manual.

**E-1**

- Optic Cover not closed. Close Cover, begin again.

**E-3**

- ANSWER strip discolored or used. Begin again with new ANSWER strip.

**E-5**

- ANSWER strip dislodged or removed before firing LANCET. Reinsert immediately.

Batteries dead or ANSWER meter malfunctioning. See Instruction Manual.

## CLEARING DISPLAY MESSAGES

1. Remove ANSWER strip.
2. Close Optic Cover.
3. Pull back Sliding Button.
4. Press Release Button.

**E-8**

Release Button not pressed within 3 minutes after insertion of ANSWER strip. Begin again with same ANSWER strip.



EP 91 30 2521

| DOCUMENTS CONSIDERED TO BE RELEVANT  |  |   |   |                 |                                  |          |           |            |                   |
|--|--|---|---|-----------------|----------------------------------|----------|-----------|------------|-------------------|
| Category   | Citation of document with indication, where appropriate, of relevant passages  | Relevant to claim   | CLASSIFICATION OF THE APPLICATION (Int. Cl.5) |                 |                                  |          |           |            |                   |
| D, X   | US-A-4 787 398 (GARCIA et al.)<br>* Abstract; column 1, line 1 - column 17, line 9; figures 1-8 *<br>-----   | 1-15  | A 61 B 5/14<br>A 61 B 5/00                    |                 |                                  |          |           |            |                   |
| X  | EP-A-0 254 203 (PERSONAL DIAGNOSTICS INC.)<br>* Abstract; column 2, line 3 - column 3, line 22; column 4, line 1 - column 6, line 48; figures 1-4 *<br>----- | 1-8, 10, 13-15  |   |                 |                                  |          |           |            |                   |
| TECHNICAL FIELDS SEARCHED (Int. Cl.5)  |  |   |   |                 |                                  |          |           |            |                   |
| A 61 B   |  |   |   |                 |                                  |          |           |            |                   |
| <p>The present search report has been drawn up for all claims</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 33%;">Place of search</td> <td style="width: 33%;">Date of completion of the search</td> <td style="width: 34%;">Examiner</td> </tr> <tr> <td>THE HAGUE</td> <td>01-07-1991</td> <td>FONTENAY P.H.E.V.</td> </tr> </table> |  |   |   | Place of search | Date of completion of the search | Examiner | THE HAGUE | 01-07-1991 | FONTENAY P.H.E.V. |
| Place of search  | Date of completion of the search   | Examiner  |   |                 |                                  |          |           |            |                   |
| THE HAGUE  | 01-07-1991   | FONTENAY P.H.E.V.   |   |                 |                                  |          |           |            |                   |
| CATEGORY OF CITED DOCUMENTS  |  | T : theory or principle underlying the invention<br>E : earlier patent document, but published on, or<br>after the filing date<br>D : document cited in the application<br>L : document cited for other reasons<br>.....<br>& : member of the same patent family, corresponding<br>document |   |                 |                                  |          |           |            |                   |
| X : particularly relevant if taken alone<br>Y : particularly relevant if combined with another<br>document of the same category<br>A : technological background<br>O : non-written disclosure<br>P : intermediate document   |  |   |   |                 |                                  |          |           |            |                   |